



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : A61K 31/495, C07D 241/12	A1	(11) International Publication Number: WO 93/23051 (43) International Publication Date: 25 November 1993 (25.11.93)
(21) International Application Number: PCT/US93/04631 (22) International Filing Date: 17 May 1993 (17.05.93) (30) Priority data: 886,021 20 May 1992 (20.05.92) US (60) Parent Application or Grant (63) Related by Continuation US 886,021 (CIP) Filed on 20 May 1992 (20.05.92) (71) Applicant (for all designated States except US): MERCK & CO., INC. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).		(72) Inventors; and (75) Inventors/Applicants (for US only) : GRAHAM, Donald, W. [US/US]; 294 Old Tote Road, Mountainside, NJ 07092 (US). ASTER, Susan, D. [US/US]; 601 Churchill Road, Teaneck, NJ 07666 (US). HAGMANN, William [US/US]; 871 Shackamaxon Drive, Westfield, NJ 07090 (US). TOLMAN, Richard, L. [US/US]; 29 Upper Warren Way, Warren, NJ 07059 (US). (74) Agent: GRASSLER, Frank, P.; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). (81) Designated States: AU, BB, BG, BR, CA, CZ, FI, HU, JP, KR, KZ, LK, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SK, UA, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: NEW 17-ESTER, AMIDE, AND KETONE DERIVATIVES OF 3-OXO-4-AZASTEROIDS AS 5A-REDUCTASE INHIBITORS (57) Abstract Described are new 17 β -ester, amide, and ketone 4-aza-5a-androstan-3-ones and related compounds and the use of such compounds as 5a-reductase inhibitors for treatment of benign prostatic hyperplasia and other hyperandrogenetic related disorders.		

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TITLE OF THE INVENTIONNEW 17-ESTER, AMIDE, AND KETONE DERIVATIVES OF 3-
OXO-4-AZASTEROIDS AS 5A-REDUCTASE INHIBITORS

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BACKGROUND OF THE INVENTION

The present invention is directed to new 17 β -ester, amide, and ketone derivatives of 4-aza-5 α -androstane-3-ones and related compounds and the use of such compounds as 5 α -reductase inhibitors.

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DESCRIPTION OF THE PRIOR ART

The art reveals that certain undesirable physiological manifestations, such as acne vulgaris, seborrhea, female hirsutism, male pattern baldness and benign prostatic hyperplasia, are the result of hyperandrogenetic stimulation caused by an excessive accumulation of testosterone or similar androgenic hormones in the metabolic system. Early attempts to provide a chemotherapeutic agent to counter the undesirable results of hyperandrogenicity resulted in the discovery of several steroidal antiandrogens having undesirable hormonal activities of their own. The estrogens, for example, not only counteract the effect of the androgens but have a feminizing effect as well. Non-steroidal antiandrogens have also been developed, for example, 4'-nitro-3'-trifluoromethyl-

isobutyranilide. See Neri, et al., Endo., Vol. 91, No. 2 (1972). However, these products, though devoid of hormonal effects, are peripherally active, competing with the natural androgens for receptor sites, and hence have a tendency to feminize a male host or the male fetus of a female host.

25

It is now known in the art that the principal mediator of androgenic activity in some target organs, e.g. the prostate, is 5 α -dihydrotestosterone, and that it is formed locally in the target organ by the action of testosterone-5 α -reductase. It is also known that inhibitors of testosterone-5 α -reductase will serve to prevent or lessen symptoms of hyperandrogenetic stimulation.

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For example, a number of 4-aza steroid compounds are known which are 5 α -reductase inhibitors. See the following Merck & Co., Inc. patents, U.S. Patent Nos. 4,377,584, 4,220,775, 4,859,681, 4,760,071 and the articles J. Med. Chem. 27, p. 1690-1701 (1984) and J. Med. Chem. 29, 2998-2315 (1986) of Rasmusson, et al., and U.S. Patent 4,845,104 to Carlin, et al., and U.S. Patent 4,732,897 to Cainelli, et al. which describe 4-aza-17 β -substituted-5 α -androstan-3-ones said to be useful in the treatment of DHT-related hyperandrogenic conditions.

Further there is the suggestion in the early prior art that hyperandrogenic diseases are the result of a single 5 α -reductase. However, there are later reports regarding the presence of other 5 α -reductase isozymes in both rats and humans. For example, in human prostate, Bruchovsky, et al. (See J. Clin. Endocrinol. Metab. 67, 806-816, 1988) and Hudson (see J. Steroid Biochem. 26, p 349-353, 1987) found different 5 α -reductase activities in the stromal and epithelial fractions. Additionally, Moore and Wilson described two distinct human reductases with peaks of activities at either pH 5.5 or pH 7-9. (See J. Biol. Chem. 251, 19, p. 5895-5900, 1976.)

Recently, Andersson and Russell isolated a cDNA which encodes a rat liver 5 α -reductase (see J. Biol. Chem. 264 pp. 16249-55 (1989). They found a single mRNA which encodes both the liver and pro-static reductases in rats. This rat gene was later used to identify a human prostatic cDNA encoding a 5 α -reductase termed "5 α -reductase 1". (See Proc. Nat'l. Acad. Sci. 87, p. 3640-3644, 1990.)

More recently, a second, human prostatic reductase (5 α -reductase 2) has been cloned with properties identified with the more abundant form found in crude human prostatic extracts. (See Nature. 354, p. 159-161, 1991.)

Further, "Syndromes of Androgen Resistance" - The Biology of Reproduction, Vol. 46, p. 168-173 (1992) by Jean O. Wilson suggests that the 5 α -reductase 1 enzyme is associated with hair follicles.

Thus, the art supports the existence of at least two genes for 5 α -reductase and two distinct isozymes of 5 α -reductase in humans. Both

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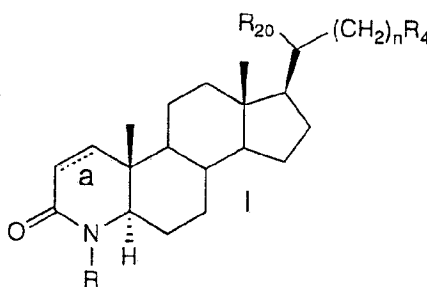
isozymes are believed to be present in prostatic tissue in which, 5 α -reductase 2, is the more abundant, while the other isozyme, 5 α -reductase 1, is believed to be more abundant in scalp tissue.

5 In the treatment of hyperandrogenetic disease conditions, e.g. benign prostatic hyperplasia (BPH) it would be desirable to have one drug entity which is dually active against both enzymes 1 and 2 in the prostate to substantially inhibit dihydrotestosterone (DHT) production. Alternatively, it would be desirable to have a drug entity which is
10 highly selective for inhibiting the scalp-associated enzyme 5 α -reductase 2. This latter drug could also be used in combination with PROSCAR® (finasteride) which is highly selective for the prostatic enzyme 5 α -reductase 2 for combination therapy in the treatment of BPH.

15 SUMMARY OF THE INVENTION

The present invention discloses novel
17 β -ester, amide, and ketone derivatives of 4-aza-5 α -androstan-3-one
compounds which are useful for inhibiting the 5 α -reductase enzyme and
isozymes thereof in prostatic tissue. They are also particularly effective
20 in selectively inhibiting the 5 α -reductase 1 associated with the scalp and/or dually inhibiting both isozymes 1 and 2 in the treatment of benign prostatic hyperplasia, acne, female hirsutism, androgenic alopecia, i.e., male pattern baldness, alopecia areata, alopecia senilis, prostatitis, and the prevention and treatment of prostatic carcinoma.

25 In accordance with the present invention there is provided novel 17 β -ester, amide, and ketone 4-aza-5 α -androstan-3-one and related compounds of the formula (I):



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wherein:

dashed line a can represent a double bond when present;

R is selected from hydrogen, methyl, ethyl, hydroxyl, amino, and methylthio;

R₂₀ is selected from hydrogen or methyl;

n is an integer from 0 to 10

R₄ is selected from:

(a) COR₁, where R₁ is C₆-C₁₀ aryl, substituted C₆-C₁₀ aryl, and heteroaryl;

(b) CONHR₂, where R₂ is substituted phenyl, heteroaryl, substituted heteroaryl, or C₇ to C₁₂ cycloalkyl;

(c) CO₂R₃, where R₃ is C₆-C₁₀ aryl, substituted C₆-C₁₀ aryl, or C₇-C₁₂ cycloalkyl;

where the above aryl and heteraryl radicals can also be fused with a benzo or another heteroaryl ring and can further be substituted with one or more substituents; and pharmaceutically acceptable salts and esters thereof.

Also disclosed are processes for their preparation, pharmaceutical formulations comprising the novel compounds as active ingredients and methods of inhibiting 5 α -reductases 1 and/or 2 in diseases which occur under hyperandrogenetic conditions, e.g., benign prostatic hyperplasia.

DETAILED DESCRIPTION OF THE

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INVENTION AND PREFERRED EMBODIMENTS

The structure I above encompasses all the 5a-reductase inhibitor compounds of this invention.

By the term C₁-C₄ alkyl is meant linear or branched alkyl; e.g. methyl, ethyl, isopropyl, propyl, n-butyl, isobutyl, sec-butyl and the like.

Dashed line "a" can independently be a double bond and when present, the compound is a delta-1-ene.

R₁ and R₃ can be a C₆-C₁₀ aryl including phenyl, benzyl, 1- and 2-phenethyl and naphthyl.

R₂ can be a phenyl group.

R₁ and R₂ can also be 5-6 membered heteroaryl radicals being fully unsaturated containing 1-4 nitrogen atoms, e.g. pyridyl, pyrrol, imidazolyl, tetrazolyl, pyrazinyl, pyrimidinyl, pyrazolyl, or triazolyl; containing 1-2 oxygen or sulfur atoms, e.g. thienyl, furanyl; or in combination with 1-2 nitrogen atoms, e.g. isothiazolyl, thiazolyl, isoxazolyl, oxazolyl or thiadiazolyl; or fused with a benzo ring, e.g. quinolyl, isoquinolyl, benzothienyl, isobenzofuryl, indolyl, carbazolyl; or fused with another heteroaryl ring, e.g. purinyl, and the like.

The C₇ - C₁₂ cycloalkyl in R₂ and R₃ can be 1-, 2-adamantyl, norbornyl, and bicyclo[2.2.2.]octyl.

The aryl or heteroaryl ring in R₁ and R₃ as well as the phenyl group in R₂ can be unsubstituted or substituted with one or more of the following substituents providing the substitution leads to a chemically inert, but biologically active 5a reductase inhibitor:

The ring substituents include:

C₁-C₈ straight or branched alkyl; e.g. methyl, ethyl, propyl, isopropyl, n-butyl, sec-butyl, isobutyl, t-butyl, n-pentyl, n-hexyl, iso-hexyl, n-butyl, n-octyl, iso-octyl, t-octyl, and the like; C₂-C₈ straight or branched alkenyl, e.g. ethenyl, 1-propenyl, 2-propenyl, 1-butenyl, 2-butenyl, 3-butenyl, 1-pentenyl, 1-hexenyl, 1-heptenyl, 1-octenyl, 2-octenyl, and the like;

C₃-C₈ cycloalkyl e.g., cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, 1-methyl-

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cyclobutyl, cyclopentyl, cyclohexyl, 1-methyl-
cyclohexyl, 2-methylcyclohexyl, 2-ethylcyclohexyl, and the like;
C₂-C₈ alkynyl e.g., 1-ethynyl; 1-propynyl, 2-propynyl, 2-

5 butynyl, 2-pentynyl, 1-hexynyl, 1-heptynyl, 1-octynyl;

CONR⁴R⁵ where R⁴ and R⁵ independently
are H, C₁-C₈ alkyl, as defined above, C₃-C₈ cycloalkyl as defined
above, C₁-C₄ perhaloalkyl

e.g., trifluoromethyl, perfluoromethyl, trichloro-
10 methyl, preferably perfluoroalkyl; phenyl, or sub-
stituted phenyl, as described below;

COR⁴, where R⁴ is defined above, including acetyl,
isobutylcarbonyl, benzoyl and the like;

S(O)_n R⁴, where n is 0-2 and R⁴ is defined above,
15 including methylsulfinyl, methylsulfonyl,
phenylsulfonyl, 4-chlorophenylsulfinyl and the like;

OCOR⁴, where R⁴ is defined above, including acetoxy,
propionyloxy, benzoyloxy, 4-chlorobenzoyloxy and the like.

SO₂NR⁴R⁵ where R⁴ and R⁵ are described above,
20 including sulfonamido, N-methylsulfonamido, N-phenylsulfonamido,
N,N-dimethylsulfonamido and the like;

NR⁴(CO)R⁵, wherein R⁴ and R⁵ are defined above,
including: acetylamino, benzoylamino, N-methylbenzoylamino and the
like;

NR⁴(CO)NHR⁵, wherein R⁴ and R⁵ are described above,
25 including: ureido, N-methylureido, N-methyl-N¹-phenylureido and the
like;

NHSO₂R⁴, R⁴ being defined above, including
methylsulfonylamino, phenylsulfonylamino and the like;

OR⁴, where R⁴ is defined above, including methoxy,
30 phenoxy, 4-chlorophenoxy and the like.

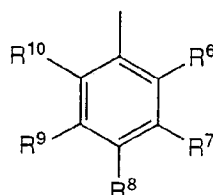
NR⁴R⁵, wherein R⁴ and R⁵ are described above, including
amino, methylamino, dimethylamino, anilino and the like;

Cyano, nitro, halo, including: fluoro, chloro, bromo and
iodo;

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Perhalo C₁-C₄ alkyl, including: tri-fluoromethyl, perfluoroethyl, trichloromethyl and the like:

CO₂R⁴, wherein R⁴ is defined above, including CO₂CH₃, CO₂Ph, CO₂-(1-adamantyl) and the like; phenyl and substituted phenyl of the formula:



wherein the radicals R⁶-R¹⁰ each can represent one or more of the substituents defined above, including; hydrogen, 4-chlorophenyl, 4-fluorophenyl, 4-methoxyphenyl, 4-phenoxy and the like.

Unless otherwise indicated the 17-substituents herein described are assumed to be in the b con-figuration

Representative compounds of the present invention include the following:

21-benzoyl-4-aza-5a-pregnan-3-one,

21-benzoyl-4-methyl-4-aza-5a-pregnan-3-one,

21-(2-methoxybenzoyl)-4-aza-5a-pregn-1-ene-3-one,

4-methyl-21-(2-trifluoromethylbenzoyl)-4-aza-5a-pregn-1-en-3-one,

20-benzoyl-4-methyl-4-aza-5a-pregnan-3-one,

23-(2-fluorobenzoyl)-4-methyl-24-nor-4-aza-5a-cholane-3,23-dione,

23-(3-pyridyl)-24-nor-4-aza-5a-chol-1-ene-3,23-dione,

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4-methyl-24-(2-pyridyl)-21-nor-4-aza-5a-cholane-3,24-dione,

5 17b-[5-(3-chlorobenzoyl)pentyl]-4-aza-5a-androstan-3-one,

17b-(6-benzoylhexyl)-4-methyl-4-aza-5a-androstan-3-one,

17b-(10-benzoyldecyl)-4-aza-5a-androst-1-en-3-one,

10

4-methyl-21-(2-thienyl)-4-aza-5a-pregnane-3,21-dione.

24-(2-pyrazinyl)-4-aza-5a-chol-1-ene-3,24-dione,

15

4-ethyl-(2,6-dimethoxybenzoyl)-4-aza-5a-pregnan-3-one,

N-(4-acetylphenyl)-4-methyl-3-oxo-4-aza-5a-pregnane-21-carboxamide,

20

N-(4-acetylphenyl)-3-oxo-4-aza-5a-pregnane-21-carboxamide,

4-methyl-3-oxo-N-(4-pyridyl)-4-aza-5a-pregnane-21-carboxamide,

25

3-oxo-N-(4-pyridyl)-4-aza-5a-pregnane-21-carboxamide,

N-(2-adamantyl)-3-oxo-4-aza-5a-pregnane-21-carboxamide,

30

N-(2-adamantyl)-4-methyl-3-oxo-4-aza-5a-pregnane-21-carboxamide,

3-oxo-N-(4-pyridyl)-4-aza-5a-pregnan-21-amide,

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- 5 4-methyl-3-oxo-4-(4-pyridyl)-4-aza-21-nor-5a-cholan-24-
amide,
- 4-methyl-3-oxo-N-(3-pyridyl)-4-aza-5a-pregnane-21-
carboxamide,
- 10 4-methyl-3-oxo-N-(2-pyridyl)-4-aza-5a-
pregnane-21-carboxamide,
- N-(1-adamantyl)-4-methyl-3-oxo-4-aza-5a-pregnane-20(S)-
carboxamide,
- 15 N-(4-acetylphenyl)-4-methyl-3-oxo-4-aza-5a-
pregnane-20(S)-carboxamide,
- N-(4-chlorophenyl)-4-methyl-3-oxo-4-aza-5a-cholan-24-
amide,
- 20 N-(4-acetylphenyl)-4-methyl-3-oxo-4-aza-5a-cholan-24-
amide,
- 3-oxo-N-(4-trifluoromethylphenyl)-4-aza-5a-
cholan-24-amide,
- 25 4-methyl-3-oxo-N-(4-pyridyl)-24-nor-4-aza-5a-
cholan-24-amide,
- N-(1-adamantyl)-11-(4-methyl-3-oxo-4-aza-5a-
androstan-17 β -yl)undecanamide,
- 30 N-(2-pyridyl)-6-(4-methyl-3-oxo-4-aza-5a-
androstan-17 β -yl)hexanamide.

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N-(3-pyridyl)-5-(3-oxo-4-aza-5a-androst-1-en-17 β -yl)pentanamide,

5 N-(2-thienyl)-7-(4-methyl-3-oxo-4-aza-5a-androstan-17 β -yl)heptanamide,

3-oxo-N-(2-pyrazinyl)-4-aza-5a-pregnan-21-amide,

10 4-methyl-3-oxo-N-(2-t-butylphenyl)-4-aza-5a-cholane-24-carboxamide,

4-methyl-3-oxo-N-(2-cyanophenyl)-4-aza-chol-1-ene-24-carboxamide,

15 N-(2-bicyclo[2.2.2]octyl)-9-(3-oxo-4-aza-5a-androstan-17 β -yl)nonanamide,

1-adamantyl 4-methyl-3-oxo-4-aza-5a-pregnane-20(S)-carboxylate,

20 phenyl 4-methyl-3-oxo-4-oxo-4-aza-5a-pregnane-20(S)-carboxylate,

25 2-(t-butyl)phenyl 4-methyl-3-oxo-4-aza-5a-pregnane-21-carboxylate,

2-methoxyphenyl 4-methyl-3-oxo-4-aza-5a-pregnane-21-carboxylate,

30 phenyl 3-oxo-4-aza-5a-pregnane-21-carboxylate,

phenyl 4-methyl-3-oxo-4-aza-5a-pregnane-21-carboxylate.

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phenyl 5-(4-methyl-3-oxo-4-aza-5a-androstan-17 β -yl)pentanoate,

- 5 2-(t-butyl)phenyl 3-oxo-4-aza-5a-pregnan-21-oate,

 2,6-dimethoxyphenyl 3-oxo-4-aza-5a-pregn-1-en-21-oate,

 2-adamantyl 8-(4-methyl-3-oxo-4-aza-5a-androstan-17 β -yl)octanoate,
10 2,6-dimethylphenyl 3-oxo-4-aza-5a-pregn-1-en-21-oate,

 2,6-dichlorophenyl 4-methyl-3-oxo-4-aza-5a-pregn-1-en-21-ate,
15 phenyl 10-(4-methyl-3-oxo-4-aza-5a-androstan-17 β -yl)decanoate,

20 and also including the corresponding compounds wherein the 4-hydrogen substituent is replaced by a methyl or an ethyl radical, and/or a delta-one double bond is present.

 Also included within the scope of this invention are pharmaceutically acceptable salts, i.e. hydrochloride, hydrobromide, acetate, pamoate, and the like, of the compound where a basic
25 heteroaryl radical is present, e.g. 4-pyridyl, which can be used as the dosage form for modifying solubility or hydrolysis characteristics or for use as sustained release or prodrug formulations.

 The novel compounds of formula I of the present invention are prepared by methods discussed below. The compounds and
30 intermediates and their physical properties are listed in Table 1, and their preparation is illustrated in Examples 1-11.

 The starting materials for these preparations are the appropriate carboxylic acids 1, 2, 9, 16, 17, 34, 37, 40, 45, 63, and 64. The source of these carboxylic acids is as follows:

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The synthesis of acid **1** is published in J.Med.Chem. **1984**, Vol. 27, p 1690.

Acid **2** was prepared by the same procedure as acid **1**.

5 The preparation of acid **9** is published in J.Med.Chem. **1984** Vol. 27, p.1690.

The syntheses of acids **16** and **17** are detailed in Example 1. The aldehyde **50** was reacted with the methyl (diethylphosphono)acetate anion to give the olefinic Horner-Wadsworth reaction product **51**. Hydrogenation to the saturated ester **52** and saponification with aqueous KOH in methanol gave **17**. The same sequence of reactions starting with the corresponding 4-methyl aldehyde (J.Med.Chem.**1986** Vol. 29, p. 2304, compound 10bg) produced the acid **16**.

15 Acid **34** was prepared as outlined in Example 10. The nitrile **58** was transformed into the methyl ester **59** by conversion to the iminoester with anhydrous HCl in methanol followed by treatment with water. KOH saponification produced the acid **34**.

20 The acid **37** was synthesized by the Arndt-Eistert homologation of **16**. Activation of **16** as the mixed anhydride with isobutyl chloroformate and N-methyl morpholine and reaction with diazomethane gave the diazoketone **53**. Silver benzoate catalyzed decomposition of **53** in methanol gave the homologous methyl ester **54**. Saponification have the corresponding acid **37**.

25 The synthesis of the acid **41** is published in J.Med.Chem. **1986** Vol. 29, p. 2300. The acids **45**, **63**, and **64** were prepared by the reaction sequence detailed in Example 11. Palladium catalyzed coupling of the D¹⁶-17-triflate **60** with methyl 4-pentynoate using the procedures published in Synlett. **1991** p. 409; J.Org.Chem. **1992** Vol. 57, p. 973. gave the enyne **61**. Hydrogenation catalyzed with palladium on carbon formed the saturated ester **62** and KOH saponification gave **45**. Similar reaction sequences using methyl 5-hexynoate and 10-undecynoic acid gave the acids **63** and **64**.

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Starting with the above acids, the novel ketones, amides, and esters listed in Table 1 were prepared using the procedures discussed below and detailed in the Examples.

5 The synthesis of the ketones 32 and 33 are given in Examples 5 and 6. Using the procedures published in J.Med.Chem. 1986 Vol. 29, p. 2310, the acids 16 and 17 were converted into the 2-thiopyridyl esters 55 and 56 by reaction with triphenylphosphine and 2,2'-dithiodipyridine.(Example 5). Low temperature reaction of these
10 2-thiopyridyl esters with phenyl-magnesium chloride produced the phenyl ketones 32 and 33 (Example 6).

The amides listed in Table 1 were prepared by a variety of procedures:

15 For the unhindered acids 16, 17, 34, 37, 41, 62, 63, and 64, 4-dimethylaminopyridine (DMAP) catalyzed carbodiimide mediated condensation with the appropriate amine or aniline was used. Either dicyclohexylcarbodiimide (Example 3) or the water soluble 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (Example 8) could be used.

20 For the more hindered acids 1 and 2, formation of the mixed anhydride with isobutyl chloroformate and N-methylmorpholine followed by reaction with the appropriate amine (Example 4) proved to be the best procedure.

25 For the most hindered acid (9), it was necessary to activate the acid for reaction with an amine by either forming the acid chloride with oxalyl chloride (Example 9) or converting it into the 2-thiopyridyl ester (Example 5).

30 With hindered or unreactive amines such as 1-adamantamine or 4-acetylaniline, reaction of the 2-thiopyridyl ester alone (Example 5) or with silver trifluoromethylsulfonate catalysis (Example 7) was used to form the amide derivatives.

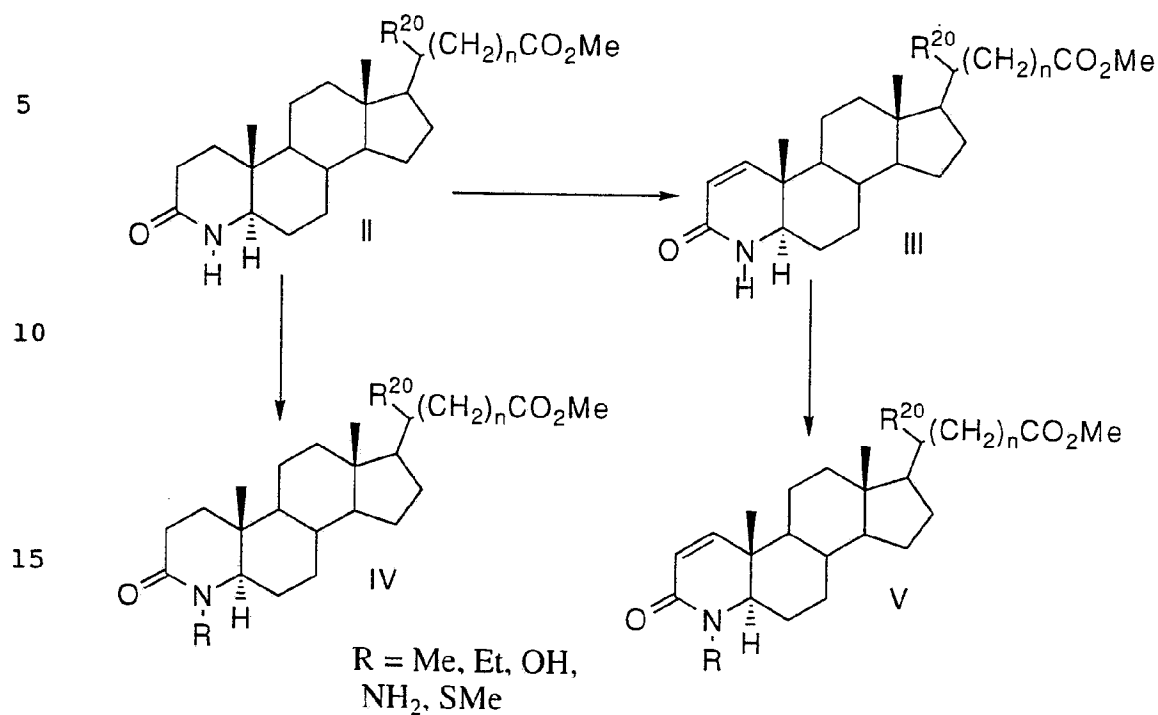
The ester derivatives in Table 1 were formed from the appropriate acid and alcohol or phenol using the same procedures (Examples 3, 7, and 8) used for the formation of the corresponding amide derivatives

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As outlined in Flowchart A the 4-H esters II can be converted into the corresponding D¹-4-H esters (III) by the procedure of Dolling, et al using dichlorodicyanobenzoquinone (J.Amer.Chem.Soc.1988, Vol. 110, p. 3318-3319) or using benzeneselenic anhydride (J.Med.Chem.1986 Vol. 29, p.2298-2315). Furthermore II and III can be alkylated on the 4-N with methyl or ethyl iodide using sodium hydride in DMF or DMSO to give the 4-methyl- or 4-ethyl-4H esters (IV) or the 4-methyl- or 4-ethyl-D¹ esters (V). Also II and III can be converted to the 4-SMe esters IV and V by reaction with sodium hydride and methanesulfonyl chloride (MeSOCl). II and III can also be aminated with hydroxylamine-O-sulfonic acid and oxidized with Oxone-acetone reagent to give the 4-NH₂ and 4-OH esters IV and V, respectively. After saponification of the esters (II-V), the starting acids containing a 4-hydrogen, methyl, ethyl, hydroxy, amino, or methylthio substituent and either with or without a D¹ can be prepared. Using the procedures in Examples 1-11, these acids can be converted into the corresponding ketone, amide, and ester derivatives.

- 15 -

Flowchart A



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The method of preparing the novel 17-ester, amide, and ketone derivatives of 3-oxo-4-azasteroids of the present invention, already described above in general terms, may be further illustrated by the following examples.

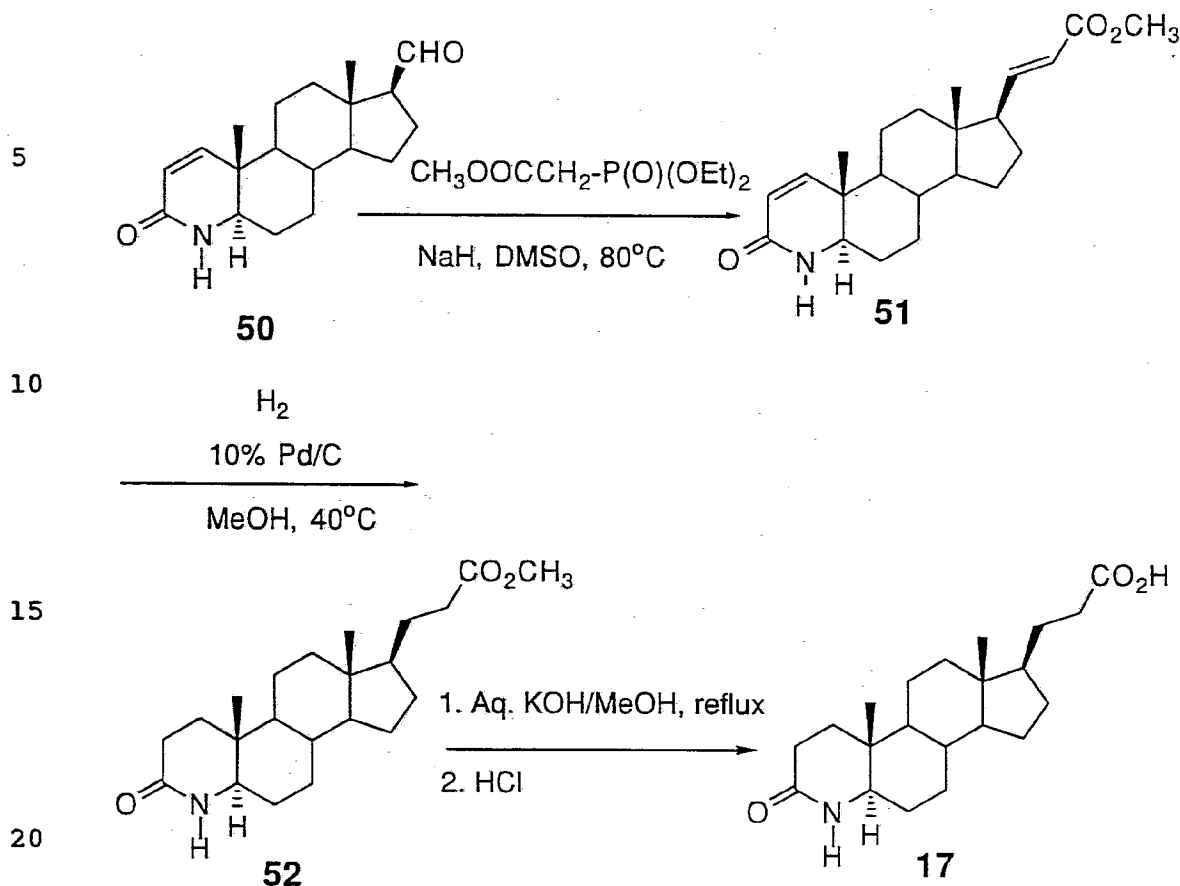
EXAMPLE 1

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3-Oxo-4-aza-5a-pregnan-21-carboxylic acid (17)

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To a suspension of the aldehyde **50**, (1.0 g, 3.32 mmole) in 12 ml DMSO was added 14.6 mg (3.65 mmole) sodium hydride (60% in mineral oil) and 698ml (3.65 mmole) methyl diethylphosphonoacetate. The mixture was heated under N₂ atmosphere at 80°C for 1 hr. The clear solution was cooled and partitioned between dilute HCl and methylene chloride. The organic phase was washed with water, brine, dried over magnesium sulfate and concentrated *in vacuo* to give 1.2 g crude product. Purification by flash chromatography on silica gel in 4:1 methylene chloride:acetone gave 956 mg of the unsaturated ester [**51**, NMR, δ =0.66(s, 3H, 18-Me), 0.97(s, 3H, 19-Me), 3.72(s, 3H, OMe), 5.7(d, 1H, D¹), 5.82(s, 1H, D²⁰), 5.96(bs, 1H, NH), 6.79(d, 1H, D¹), 6.92(dd, 1H, D²⁰)].

The unsaturated ester (**51**) (956 mg, 2.67 mmole) was dissolved in 80 ml warm methanol and hydrogenated with 300 mg 10%

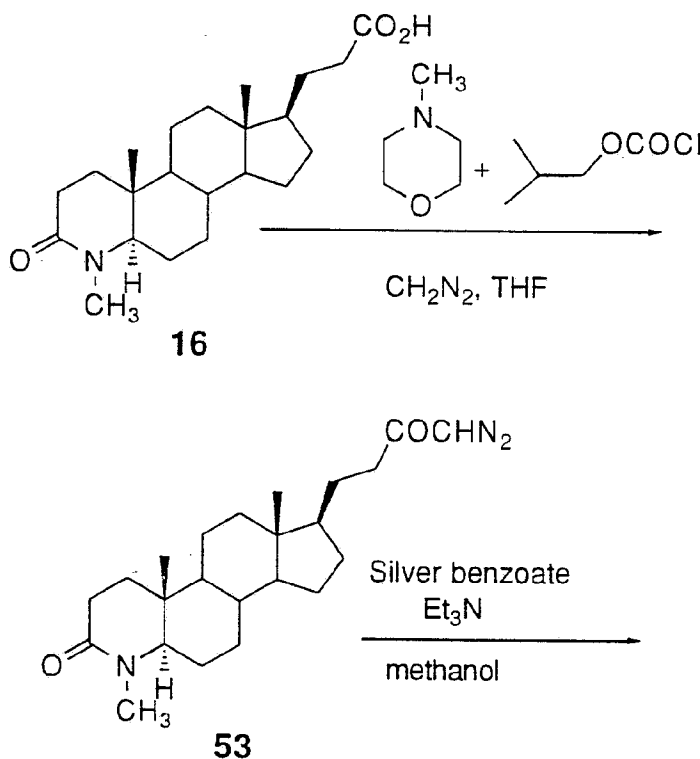
- 17 -

Pd/C at 40psi at a temperature of 40°C for 3 hrs. The mixture was filtered through a pad of Celite, washing with warm methanol. The filtrate was concentrated *in vacuo* to give 938 mg of the saturated ester (52).

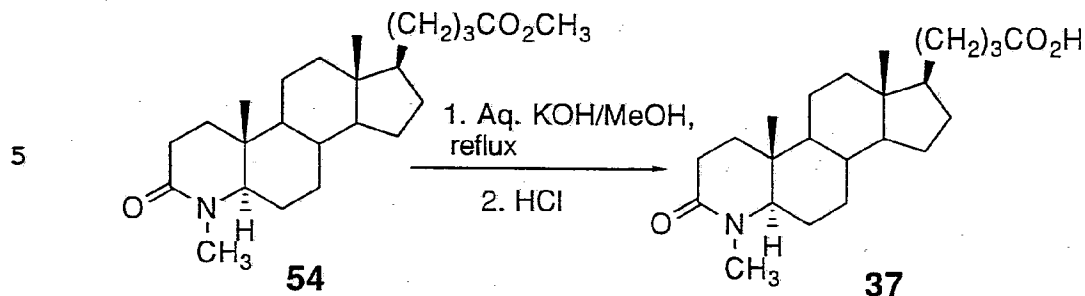
The saturated ester (52) (938 mg, 2.59 mmole) was dissolved in 9 ml methanol containing 863 ml 9M KOH and refluxed for 1 hr. The mixture was concentrated to a small volume and 100 ml water added. The mixture was cooled to 10°C and brought to pH 1 with concentrated HCl. The resulting precipitate was filtered, washed with water, sucked dry and dried in a vacuum oven at 60°C, 25 in. for 18 hrs, giving 830 mg of the acid (17).

EXAMPLE 2

3-Oxo-4-methyl-4-aza-5a-21-norcholanic acid (37)



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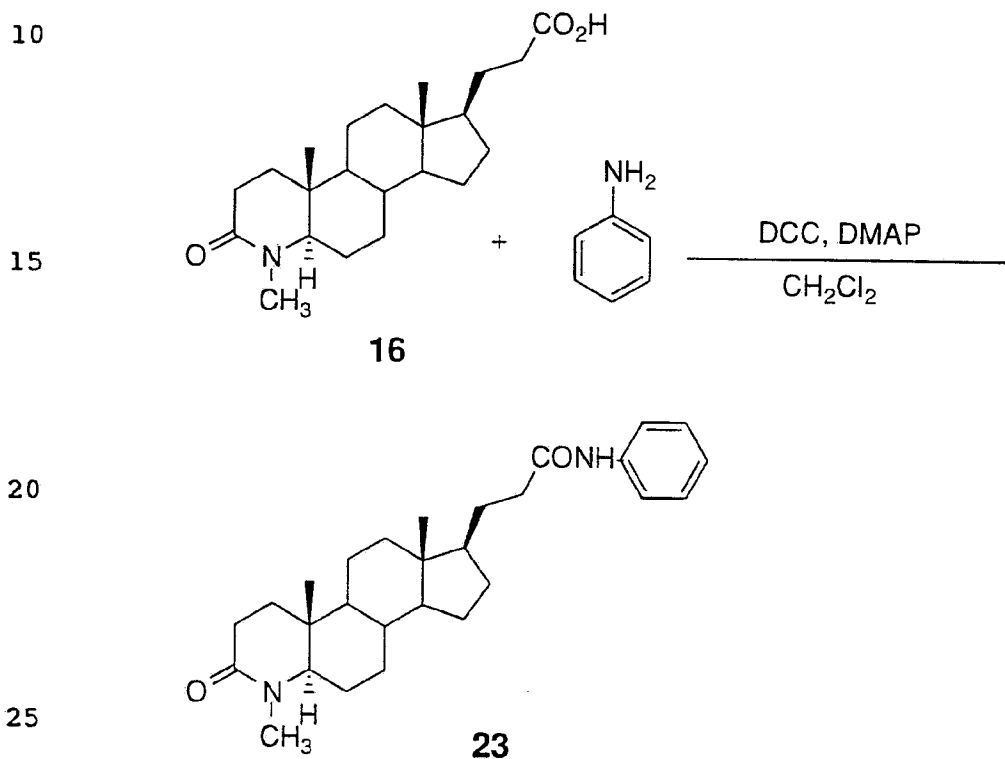
10 To a solution of the acid (**16**), (763 mg, 2.66 mmole) and N-methyl morpholine (296 ml, 2.66 mmole) in 60 ml THF at -20°C under N₂ atmosphere was added dropwise isobutyl chloroformate (352 ml, 2.66 mmole). The mixture was stirred at -20°C for 40 minutes, filtered and concentrated *in vacuo* to about 1/4 its volume on a rotary evaporator using no heat. The mixture was cooled to -10°C under N₂ and a freshly prepared ether solution of excess (12 mmole) diazomethane added. After stirring for 18 hours at room temperature, nitrogen was bubbled through the solution for 20 minutes (to remove excess diazomethane) and the mixture concentrated *in vacuo*. The residue was partitioned with methylene chloride-water and the organic phase washed with 5% acetic acid, water, brine, dried over magnesium sulfate and concentrated *in vacuo* to give 1.1 g crude product (oil). Purification by flash chromatography on silica gel in 4:1 methylene chloride:acetone gave 350 mg of the diazoketone (**53**).

20 The diazoketone (**53**), (224 mg, 0.681 mmole) was dissolved in 1 ml of methanol and 266 ml of a 0.218M solution of silver benzoate in triethyl amine (50 mg/ml) was added. The mixture was stirred at room temperature for 2 hours and the dark solution concentrated *in vacuo*. Methylene chloride was added and the mixture filtered. The filtrate was washed with dilute HCl, water, saturated NaHCO₃, brine, dried over magnesium sulfate and concentrated *in vacuo* to give 212 mg crude product (oil). Purification by flash chromatography on silica gel in 4:1 methylene chloride:acetone gave 153 mg of the methyl butyrate (**54**).

- 19 -

The methyl ester (**54**) was converted to the free acid (**37**) using the same conditions as was used in the previous case (**52** converted to **17**) except that the reflux time was 2 hours.

5

EXAMPLE 33-Oxo-4-methyl-N-phenyl-4-aza-5 α -pregnan-21-carboxamide (**23**)

To a solution of the acid (**16**), (57 mg, 0.158 mmole) in 2 ml methylene chloride was added 17 ml (0.190 mmole) aniline, 45 mg (0.220 mmole) dicyclohexylcarbodiimide (DCC) and 1 mg 4-dimethylaminopyridine (DMAP). The mixture was stirred at room temperature for 5 hours, filtered and concentrated *in vacuo* to give 96 mg crude mixture. Purification by preparative thin layer chromatography on a 1500 m silica gel plate in 5% methanol/methylene chloride and trituration with hexane gave 26 mg of the anilide (**23**).

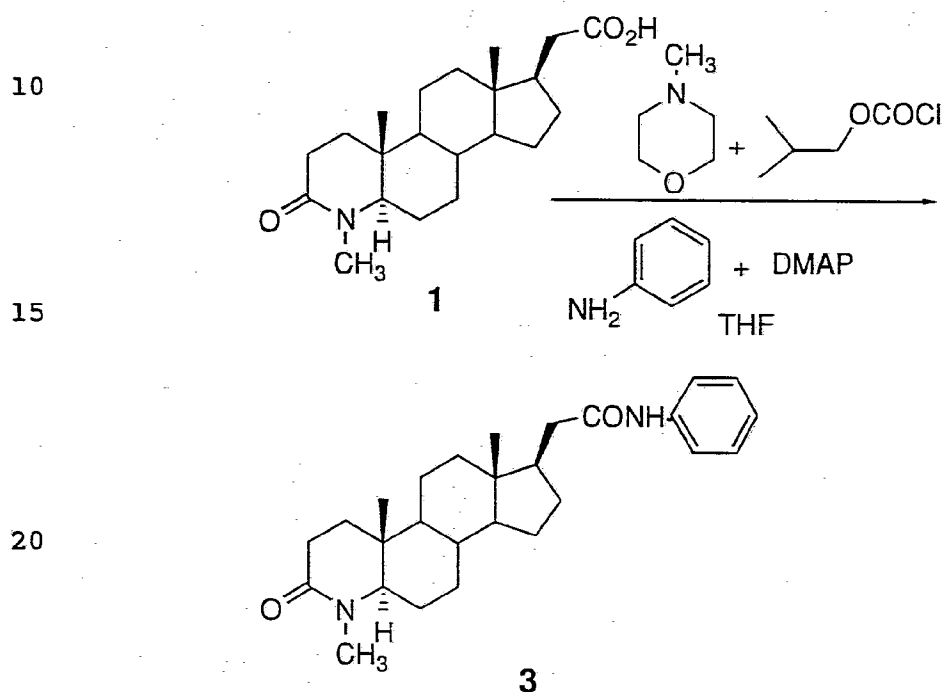
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- 20 -

Compounds 18-20, 24-28, 30, 38-39, 42 and 44 were prepared by the above procedure.

EXAMPLE 4

3-Oxo-4-methyl-N-phenyl-4-aza-5a-pregnan-21-amide (3)



25 To a solution of the acid (1) (174 mg, 0.50 mmole) and N-methyl morpholine (61 ml, 0.55 mmole) in 10 ml THF at -20°C under a N₂ atmosphere was added dropwise isobutyl chloroformate (66 ml, 0.50 mmole). The mixture was stirred at -20°C for 20 minutes and aniline (64 ml, 0.70 mmoles) plus 4-dimethylaminopyridine (DMAP, 3 mg) was added. The mixture was stirred at -20°C for 30 minutes and then

30 at room temperature for 18 hours. The mixture was concentrated *in vacuo* and the residue partitioned with methylene chloride-water. The organic phase was washed with dilute HCl, water, brine, dried over magnesium sulfate and concentrated *in vacuo* to give 224 mg crude

- 21 -

product. Purification by flash chromatography on silica gel in 4:1 methylene chloride:acetone gave 184 mg of the anilide (3).

5 Compounds **4-8**, **29** and **31** were prepared by the above procedure.

EXAMPLE 5

10 N-2-adamantyl-3-oxo-4-methyl-4-aza-5a-pregnan-21-carboxamide (21)

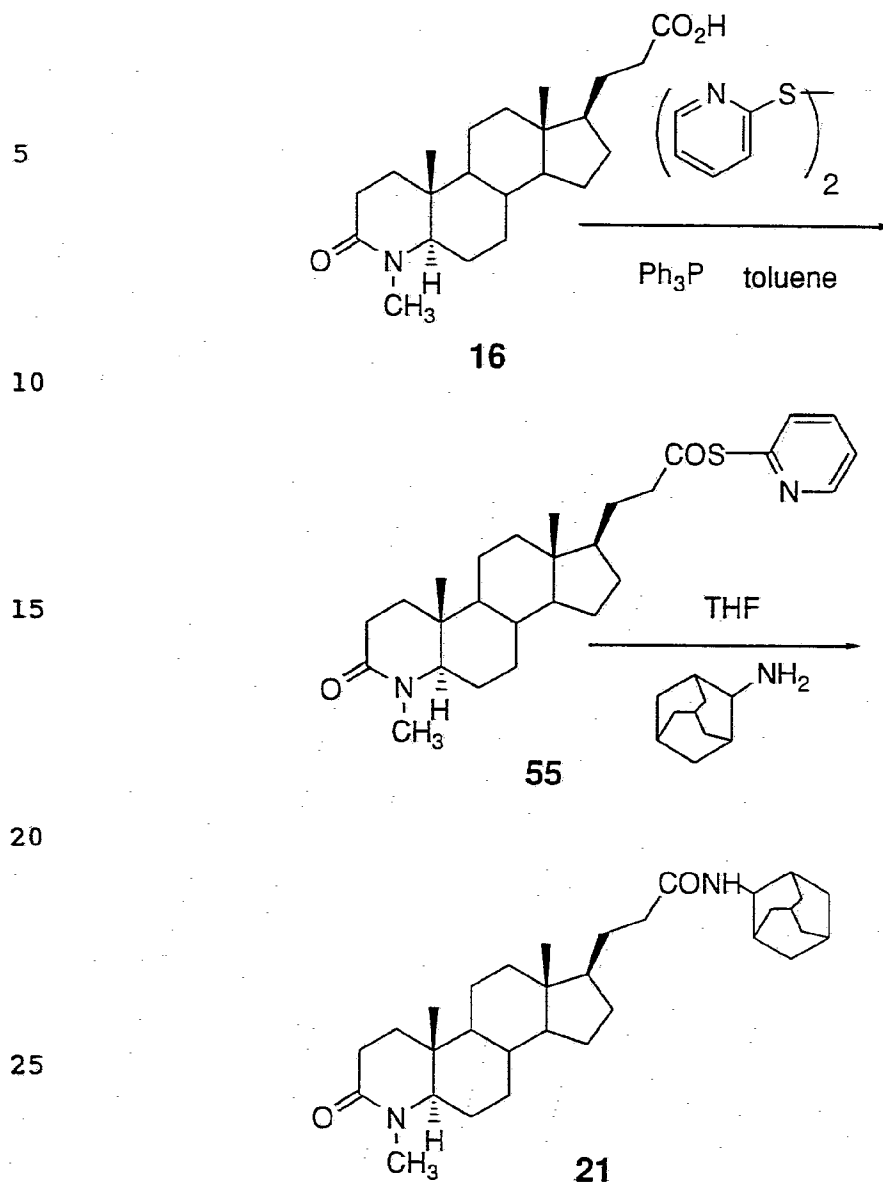
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- 22 -



To a suspension of the acid (**16**), (391 mg, 1.08 mmole) in 3 ml toluene was added 487 mg (2.16 mmole) 2,2'-dithiopyridine followed by 567 mg (2.16 mmole) triphenylphosphine. The mixture was stirred at room temperature for 18 hours, concentrated *in vacuo* and the residue flash chromatographed on silica gel in 4:1 methylene chloride:acetone to give a yellow solid. The solid was washed with ether to give 326 mg of the thiopyridyl ester (**55**).

- 23 -

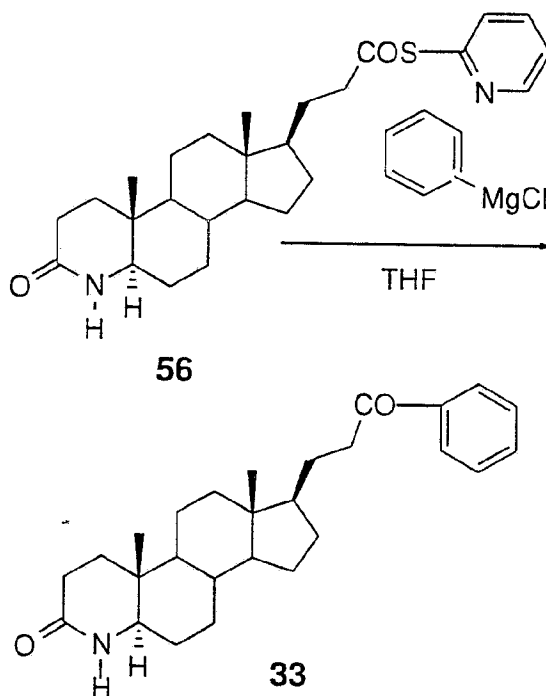
Using the same procedure, the thiopyridyl esters **56** and **57** were prepared from the carboxylic acids **17** and **9** respectively.

To a solution of the thiopyridyl ester (**55**), (105 mg, 0.231 mmole) in 2.5 ml THF was added 262 mg (1.73 mmole) 2-adamantane amine. The mixture was stirred at room temperature for 18 hours, concentrated *in vacuo* and the residue partitioned with methylene chloride-2N HCl. The organic phase was washed with water, brine, dried over magnesium sulfate and concentrated *in vacuo*. The residue was flash chromatographed on silica gel in 4:1 methylene chloride:acetone to give 83 mg of the amide (**21**).

Using the above procedure compounds **14** and **22** were prepared from the thiopyridyl esters **57** and **56** respectively.

EXAMPLE 6

21- Benzoyl-4-aza-5a-pregnan-3-one (33)



- 24 -

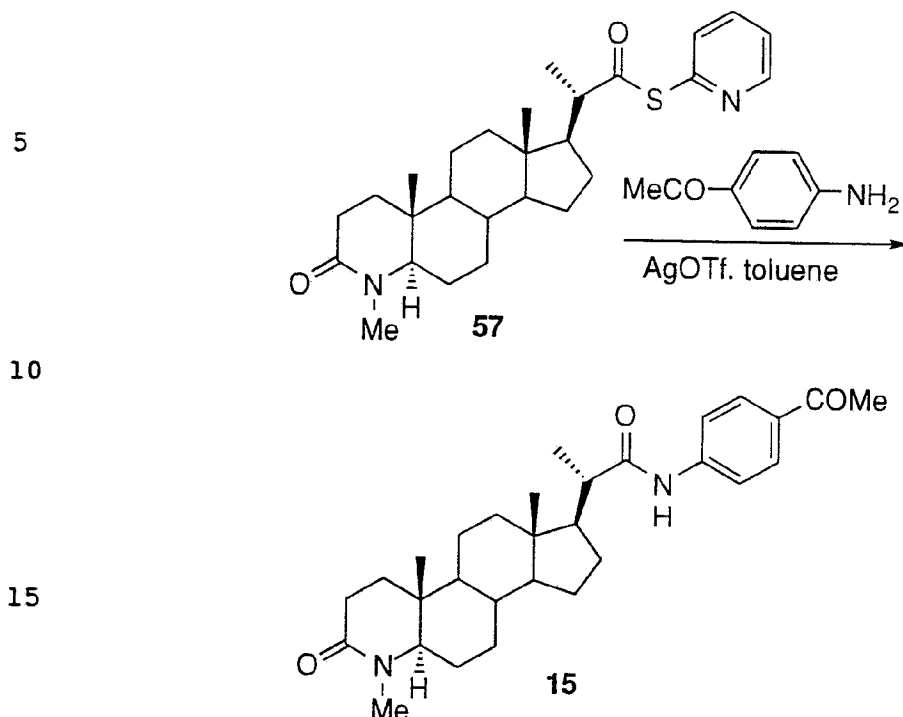
To a suspension of the thiopyridyl ester (**56**), (100 mg, 0.227 mmole) in 4 ml THF at -78°C under a N₂ atmosphere was added phenyl magnesium chloride (250 ml, 0.499 mmole, 2M in THF). The mixture was stirred at -78°C for 45 minutes and allowed to warm to 0°C. The reaction was quenched by the careful addition of 5 drops of brine. The mixture filtered and washed with THF and methylene chloride. The filtrate was concentrated *in vacuo* and the residue dissolved in methylene chloride, washed with 2N NaOH, water, brine, dried over magnesium sulfate and concentrated *in vacuo* to give 84 mg crude mixture. Purification by preparative thin layer chromatography on a 2000 m silica gel plate in 4:1 methylene chloride:acetone (run up the plate 4 times) gave 20 mg of the phenyl ketone (**33**).

Compound **32** was prepared by the above procedure.

EXAMPLE 7

N-(4-Acetylphenyl)-4-methyl-3-oxo-4-aza-5a-pregnane-20(S)-carboxamide (15)

- 25 -



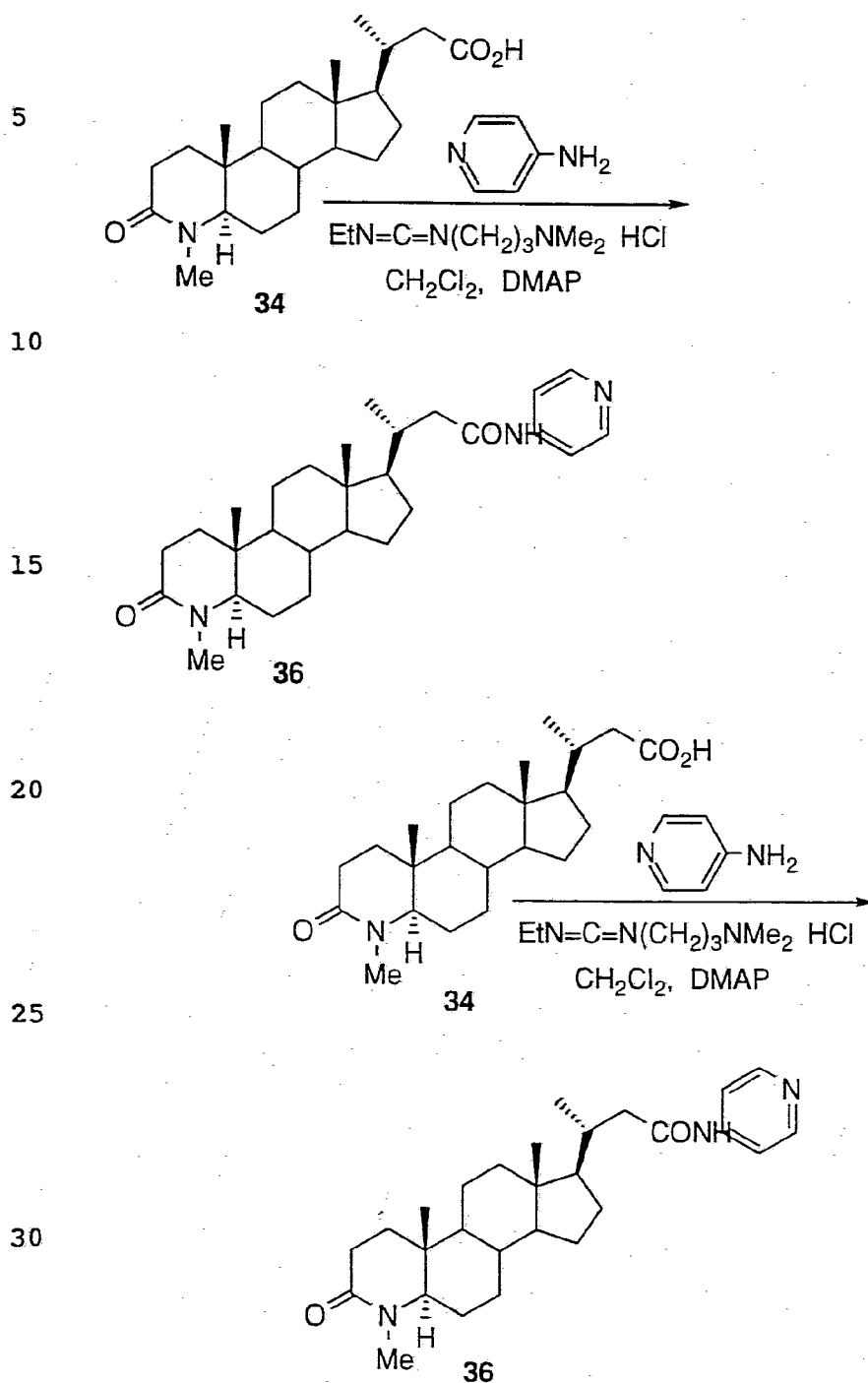
20 To a suspension of the thiopyridyl ester (**57**) (136 mg, 0.3 mmoles) and 4-acetylaniline (81 mg, 0.6 mmoles) in 1.5 ml of toluene was added all at once a warm solution of 93 mg (0.36 mmoles) of silver trifluoromethanesulfonate in 0.6 ml of toluene. The suspension was stirred at room temperature for 24 hrs. Periodically the gummy precipitate was dispersed with a glass rod. The suspension was diluted with 30 ml of CH₂Cl₂ and washed with water, 5% ammonium hydroxide, water, and saturated brine and dried (MgSO₄). Evaporation *in vacuo* gave 117 mg of a pale yellow solid. Preparative TLC on two 20x20 cm, 1000 m, silica gel plates with 4:1 CH₂Cl₂-acetone and elution of the strongly UV-active band with 4:1 CH₂Cl₂-MeOH gave 32 mg of pure amide **15**.

30 Compounds **10** and **11** were prepared by the above procedure.

EXAMPLE 8

4-Methyl-3-oxo-N-(4-pyridyl)-24-nor-4-aza-5 α -cholan-23-amide(**36**)

- 26 -



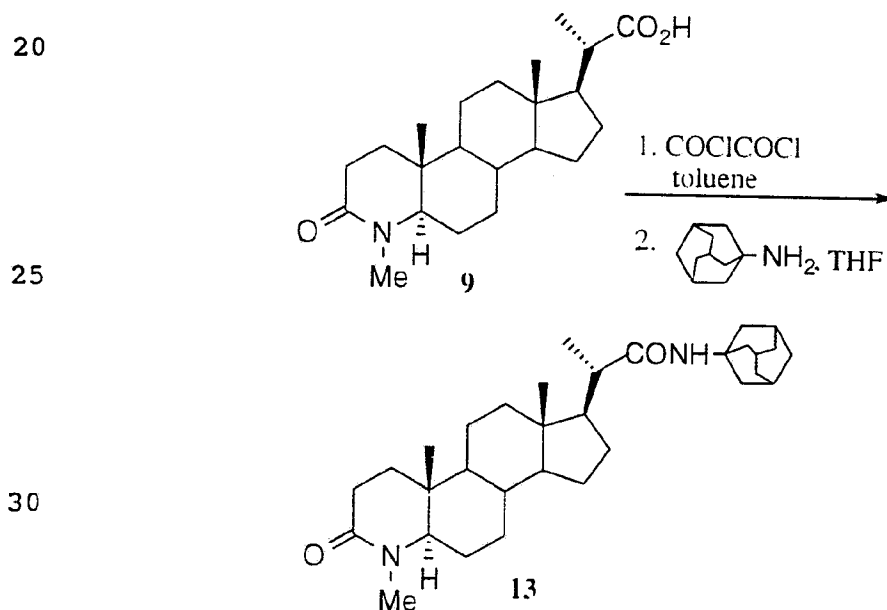
- 27 -

To a solution of 38 mg (0.1 mmole) of the acid **34**, 19 mg (0.2 mmole) of 4-aminopyridine, and 1 mg of 4-dimethylaminopyridine in 0.3 ml of CH₂Cl₂ was added 38 mg (0.2 mmole) of 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride. The solution was kept at room temperature for 24 hrs, diluted with CH₂Cl₂, washed with water, dilute K₂CO₃, water, and saturated brine and dried (MgSO₄). Evaporation *in vacuo* gave 45 mg of a gum. Preparative TLC on a 20x20 cm, 1000 m silica gel plate with 7% MeOH in CH₂Cl₂ and elution of the strongly UV- active band with 2:1 CH₂Cl₂-MeOH gave 23 mg of pure amide **36**.

Compounds **35**, **43**, **46**, **47**, **48**, and **49** were prepared from the appropriate acids by the above procedure.

EXAMPLE 9

N-(1-Adamantyl)-4-methyl-3-oxo-4-aza-5a-pregnane-20(S)-carboxamide(13).



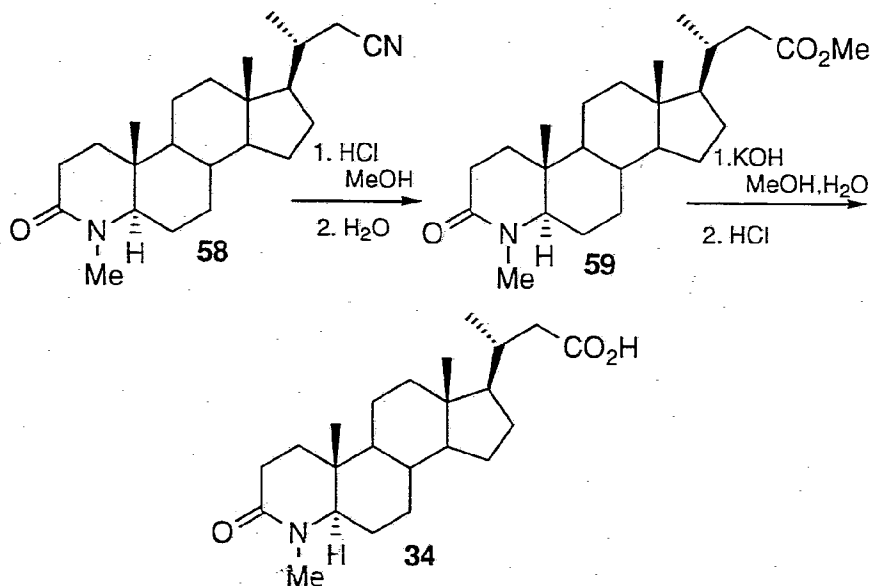
To a suspension of 108 mg (0.3 mmole) of the acid **9** in 1.3 ml of toluene was added 0.24 ml (2.75 mmole) of oxalyl chloride

- 28 -

at room temperature. After 30 min the volatiles were removed *in vacuo*, and the residue was dissolved in 1.5 ml of THF. A solution of 181 mg (1.2 mmoles) of 1-adamantanamine and 3 mg of 4-dimethylaminopyridine in 1 ml of THF was added, and the mixture allowed to stir at room temperature for 20 hrs. Most of the THF was removed *in vacuo*, and the residue partitioned between ice water and CH₂Cl₂. The aqueous phase was extracted with CH₂Cl₂ (2X). The combined organic extracts were washed with 0.5 N HCl, water, and dried (MgSO₄). Evaporation *in vacuo* and flash chromatography of the residue on a 10 mm x 18 cm column of silica gel with 1:1 ethyl acetate-acetone gave 47 mg of pure amide 13.

EXAMPLE 10

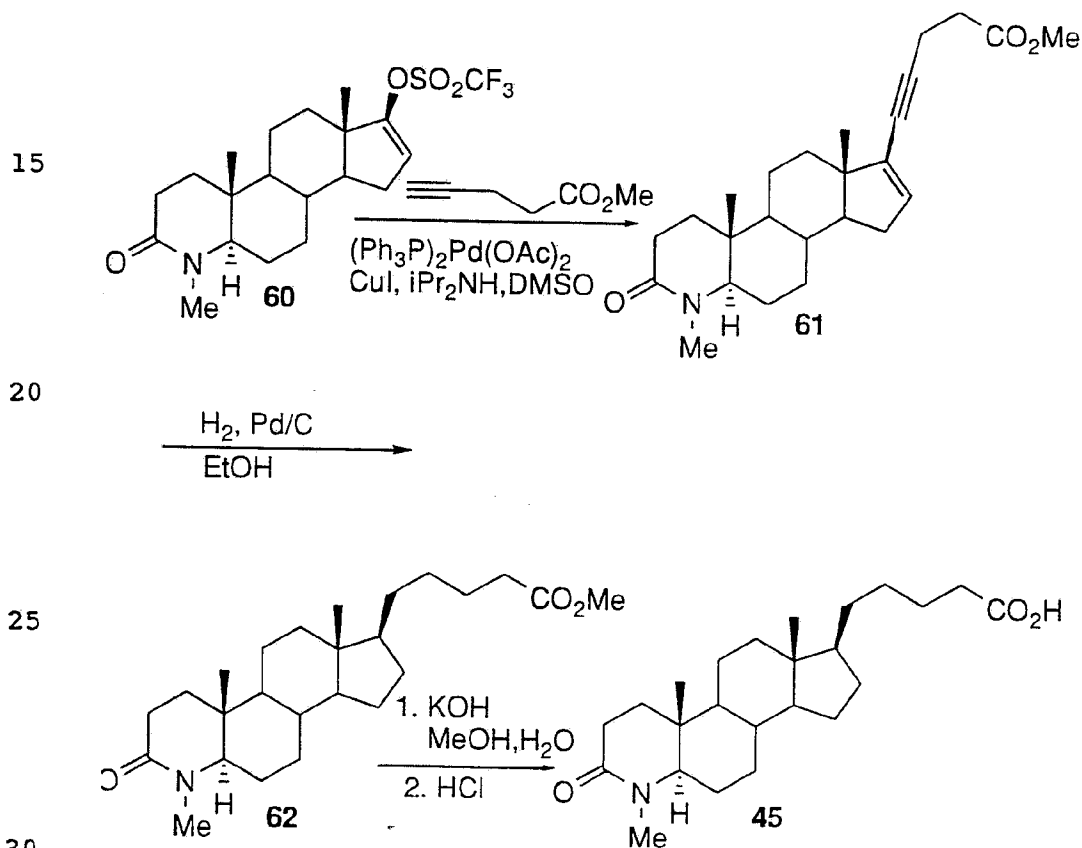
4-Methyl-3-oxo-24-nor-4-aza-5a-cholan-23-oic acid(34)



A solution of 144 mg of 4-methyl-3-oxo-24-nor-4-aza-5a-cholano-23-nitrile (58) in 5 ml of methanol saturated with anhydrous HCl was allowed to stand at room temperature for 6 hrs. The residue after evaporation *in vacuo* was stirred in 10 ml of water for 3 hrs and extracted with CH₂Cl₂ (3X). The extracts were washed with water and

- 29 -

dried (MgSO₄). Evaporation *in vacuo* and flash chromatography of the residue on silica gel with 4:1 CH₂Cl₂-acetone gave 112 mg of methyl 4-methyl-3-oxo-24-nor-4-aza-5a-cholan-23-oate (**59**). The methyl ester **59** (101 mg) was saponified to 90 mg of the acid **34** using the procedure in Example 1.

EXAMPLE 1110 4-Methyl-3-oxo-21-nor-4-aza-5a-cholane-24-carboxylic acid(**45**)

A mixture of 436 mg (1.0 mmole) of 4-methyl-17-trifluoromethylsulfonyloxy-4-aza-5a-androst-16-en-3-one (**60**) 168 mg (1.5 mmole) of methyl 4-pentynoate, 40 mg of bis(triphenylphosphine)palladium(II) acetate, 5 mg of cuprous iodide in 3.0 ml of DMSO and 3.0 ml of N,N-diisopropylamine was stirred at

- 30 -

room temperature for 17 hrs. The dark reaction mixture was poured into 50 ml of 0.5 M HCl and extracted with CH₂Cl₂ (3X). The combined extracts were washed with water (4X) and dried (MgSO₄).

5 Evaporation *in vacuo* and flash chromatography of the residue on silica gel with 6:1 CH₂Cl₂-acetone gave 400 mg of enyne **61**. The enyne was immediately hydrogenated in 20 ml of EtOH with 150 mg of 10% palladium on carbon catalyst under a hydrogen-filled balloon. After stirring at room temperature for 24 hrs, the reaction mixture was
10 filtered through a bed of Celite., which was washed with EtOH (4X). The filtrate and washes were evaporated *in vacuo* and the residue flash chromatographed on silica gel with 5:1 CH₂Cl₂ to give 345 mg of methyl 4-methyl-3-oxo-21-nor-4-aza-5a-cholane-24-carboxylate (**62**).

The methyl ester **62** (300 mg) was saponified to 275 mg of the acid
15 **45** using the procedure in Example 1.

The acids 6-(4-methyl-3-oxo-4-aza-5a-androstan-17B-yl)hexanoic acid (**63**) and 11-(4-methyl-3-oxo-4-aza-5a-androstan-17B-yl)-undecanoic acid (**64**) used for the preparation of the anilides **48** and **49** respectively were prepared by the above procedure.

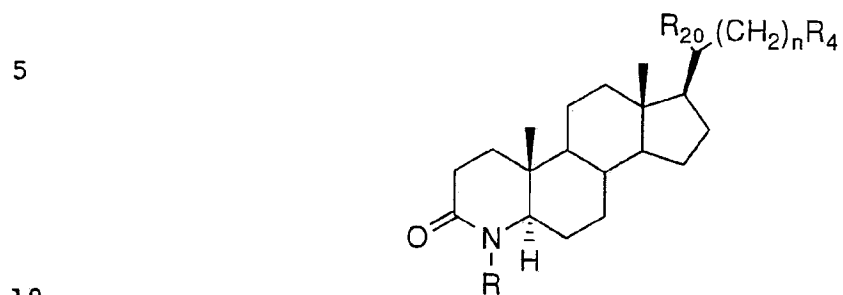
20 The following table lists the compounds made in this invention and their physical properties.



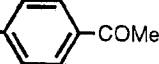
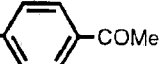



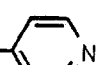
The following tables 1 and 2 list some of the compounds made in this invention and their physical properties. The compound numbers in these two tables are with reference to compounds described in Examples 1
25 through 11.

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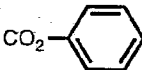
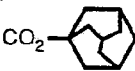
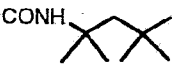
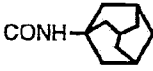
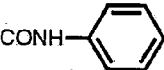

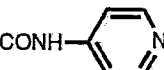
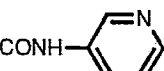
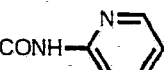


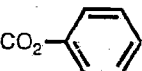
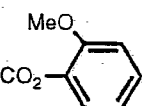
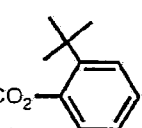
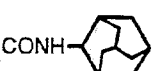
- 31 -

Table 6




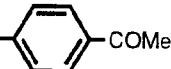

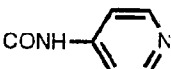
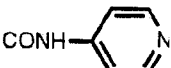
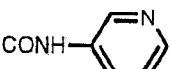
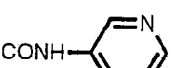
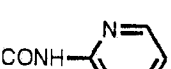
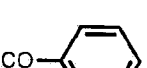
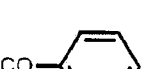
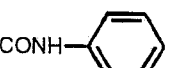
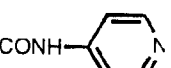


<u>Compd</u>	<u>R</u>	<u>R₂₀</u>	<u>n</u>	<u>R₄</u>	<u>NMR</u>
					18-Me
15	1	Me	H	0	CO ₂ H
	2	H	H	0	CO ₂ H
	3	Me	H	0	CONH-  0.62
20	4	H	H	0	CONH-  0.65
	5	H	H	0	CONH-  -COMe 0.66
	6	H	H	0	CONH-  -COMe 0.66
25	7	Me	H	0	CONH-  0.64
	7a	Me	H	0	CONH(Me)-  0.44
	8	H	H	0	CONH-  0.62
30	8a	H	H	0	CONH(Me)-  0.45
	9	Me	Me	0	CO ₂ H





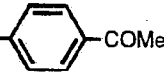







- 32 -

5	10	Me	Me	0		0.76
	11	Me	Me	0		0.67
	12	Me	Me	0		0.68
	13	Me	Me	0		0.68
10	14	Me	Me	0		0.74
	15	Me	Me	0		0.72
	15a	Me	Me	0		0.71
	15b	Me	Me	0		0.72
20	15c	Me	Me	0		0.71
	16	Me	H	1		0.60
	17	H	H	1		
	18	Me	H	1		0.65
25	19	Me	H	1		0.64
	20	Me	H	1		0.65
	21	Me	H	1		0.62

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5	22	H	H	1	CONH- 	0.62
	23	Me	H	1	CONH- 	0.63
	24	H	H	1	CONH- 	0.63
	25	Me	H	1	CONH- 	0.63
10	26	H	H	1	CONH- 	0.63
	27	Me	H	1	CONH- 	0.64
	28	H	H	1	CONH- 	0.63
	29	Me	H	1	CONH- 	0.63
20	30	H	H	1	CONH- 	0.62
	31	Me	H	1	CONH- 	0.62
	32	Me	H	1	CO- 	0.64
	33	H	H	1	CO- 	0.63
30	34	Me	Me	1	CO ₂ H	0.69
	35	Me	Me	1	CONH- 	0.73
	36	Me	Me	1	CONH- 	0.71

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	37	Me	H	2	CO ₂ H	0.58
	38	Me	H	2	CONH- 	
5	39	Me	H	2	CONH- 	0.59
	40	Me	Me	2	CO ₂ H	
	41	H	Me	2	CO ₂ H	0.65
10	42	Me	Me	2	CONH- 	0.68
	43	H	Me	2	CONH- 	0.69
15	44	Me	Me	2	CONH- 	0.68
	44a	Me	Me	2	CONH- 	0.66
	44b	Me	Me	2	CONH- 	0.66
20	44c	Me	Me	2	CONH- 	0.67
	45	Me	H	3	CO ₂ H	0.55
25	46	Me	H	3	CO ₂ - 	0.57
	47	Me	H	3	CONH- 	0.56
	48	Me	H	4	CONH- 	0.55
30	49	Me	H	9	CONH- 	0.56
	52	H	H	1	CO ₂ Me	0.61

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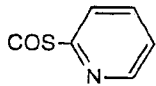
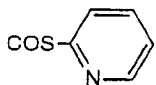
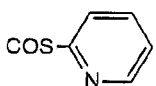
5	53	Me	H	1	COCHN ₂	
	54	Me	H	2	CO ₂ Me	0.58
	55	Me	H	1		0.62
	56	H	H	1		0.64
	57	Me	Me	0		0.74
10	58	Me	Me	1	CN	
	59	Me	Me	1	CO ₂ Me	0.70
	59a	H	Me	2	CO ₂ Me	0.64
	62	Me	H	3	CO ₂ Me	0.56
	64	Me	H	9	CO ₂ H	0.56

Table 6. part 2

	<u>Compd</u>	<u>NMR</u>	<u>Other</u>	<u>Mass Spectrum</u>
		<u>19-Me</u>		
25	1	<u>J. Med. Chem. 1984, 27, 1690-1701.</u>		
	2			m/e334(M+1) FAB
	3	0.84	7.04-7.52 (m, 5H, ArH)	m/e423(M+1) FAB
	4	0.91	7.07-7.53 (m, 5H, ArH)	m/e409(M+1) FAB
30	5	0.90	2.58(s, 3H, COMe); 7.61(d, 2H, ArH); 7.94(d, 2H, ArH)	m/e466(M+2) FAB
	6	0.90	2.57(s, 3H, COMe); 7.62(d, 2H, ArH); 7.93(d, 2H, ArH)	m/e451(M+1) FAB

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	7	0.90	7.56(d,2H,ArH); 8.46(d,2H,ArH)	m/e424(M+1) FAB
5	7a	0.85	3.30(s,3H,NMe); 7.13(d,2H,ArH); 8.65(d,2H,ArH)	
	8	0.88	7.58(d,2H,ArH); 8.44(d,2H,ArH)	m/e410(M+1) FAB
	8a	0.87	3.27(s,3H,NMe); 7.15(d,2H,ArH); 8.66(d,2H,ArH)	m/e423(M) EI
10	9		J. Med. Chem. 1984, 27 1690-1701	
	10	0.91	1.36(s,3H,21-Me); 7.0-7.5(m,5H,ArH)	
	11	0.88	1.13(d,3H,21-Me); 1.64,2.09,2.14(bs,15H,adamantylH);	
15	12	0.90	1.03(s,9H,CMe3); 1.12(d,3H,21-Me); 1.40,1.42(s,6H,NCMe2); 5.20(s,1H,NH)	
	13	0.89	1.14(d,3H,21-Me); 1.67,1.99,2.06(bs,15H,adamantylH); 5.03(bs,1H,NH)	
20	14	0.90	1.28(d,3H,21-Me); 7.06- 7.70(m,5H,ArH)	
	15	0.89	1.28(d,3H,21-Me); 2.58(s,3H,COMe); 7.68(d,2H,ArH); 7.92(d,2H,ArH); 8.00(bs,1H,NH)	
25	15a	0.89	1.27(d,3H,21-Me); 7.64(d,2H,ArH); 8.44(d,2H,ArH); 8.50(s,1H,NH)	
	15b	0.89	1.29(d,3H,21-Me); 7.28(m,1H,ArH); 7.92(s,1H,ArH); 8.30(m,2H,ArH); 8.61(bs,1H,NH)	
30	15c	0.88	1.28(d,3H,21-Me); 7.04(m,1H,ArH); 7.78(m,1H,ArH); 8.13(m,3H,ArH+NH)	
	16	0.90		m/e361(M) EI
	17			m/e347(M) EI

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	18	0.90	7.05-7.41(m,5H,ArH)	m/e437(M)	EI
	19	0.89	3.81(s,3H,OMe); 6.90-7.29(m,4H,ArH)		
5	20	0.90	1.35(s,9H,CMe ₃); 6.95-7.43(m,4H,ArH)		
	21	0.90		m/e495(M+1)	FAB
10	22	0.91		m/e481(M+1)	FAB
	23	0.89	7.05-7.54(m,5H,ArH)	m/e436(M)	EI
	24	0.91	7.06-7.55(m,5H,ArH)	m/e422(M)	EI
15	25	0.89	2.56(s,3H,COMe); 7.64(d,2H,ArH); 7.93(d,2H,ArH)	m/e478(M)	EI
	26	0.91	2.57(s,3H,COMe); 7.63(d,2H,ArH); 7.91(d,2H,ArH)	m/e464(M)	EI
	27	0.90	7.84(d,2H,ArH); 8.43(d,2H,ArH)	m/e437(M)	EI
20	28	0.91	7.58(d,2H,ArH); 8.46(d,2H,ArH)	m/e424(M+1)	FAB
	29	0.89	7.30(m,1H,ArH); 8.28(m,2H,ArH); 8.60(s,1H,ArH)		
25	30	0.90	7.33(m,1H,ArH); 8.32(m,2H,ArH); 8.62(s,1H,ArH)	m/e424(M+1)	FAB
	31	0.89	7.05(m,1H,ArH); 7.31(m,1H,ArH); 8.24(m,2H,ArH)		
	32	0.91	7.38-7.62(m,3H,ArH); 7.95(d,2H,ArH)		
30	33	0.90	7.05-7.54(m,5H,ArH)	m/e408(M+1)	FAB
	34	0.86	1.01(d,2H,21-Me)		
	35	0.89	1.04(d,3H,21-Me); 7.03-7.62(m,5H,ArH); 7.21(bs,1H,NH)		

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	36	0.87	1.01(d,3H,21-Me); 7.60(bs,2H,ArH); 8.40(s,1H,NH); 8.44(bs,2H,ArH)	
	37	0.90		m/e376(M+1) FAB
5	38	0.87	7.07-7.49(m,5H,ArH)	m/e451(M+1) FAB
	39	0.89	7.58(d,2H,ArH); 8.46(d,2H,ArH)	m/e451(M+1) FAB
	40		J. Med. Chem. 1986, 29 2298-2315	
10	41	0.88	0.91(d,3H,21-Me)	
	42	0.89	0.96(d,3H,21-Me); 7.03- 7.56(m,5H,ArH); 7.48(bs,1H,NH)	
	43	0.91	0.96(d,3H,21-Me); 5.92(bs,1H,NH); 7.04-7.64(m,5H,ArH)	
15	44	0.89	0.96(d,3H,21-Me); 2.57(s,3H,COMe); 7.66(d,2H,ArH); 7.90(d,2H,ArH); 8.16(bs,1H,NH)	
	44a	0.88	0.94(d,3H,21-Me); 7.60(d,2H,ArH); 8.37(s,1H,NH); 8.44(d,2H,ArH)	
	44b	0.88	0.96(d,3H,21-Me); 7.33(m,1H,ArH); 8.02(bs,1H,ArH); 8.33(m,2H,ArH); 8.64(s,1H,NH)	
20	44c	0.87	0.95(d,3H,21-Me); 7.08(m,1H,ArH); 7.78(m,1H,ArH); 8.25(m,2H,ArH); 8.68(bs,1H,NH)	
	45	0.87	2.31(t,2H,CH ₂ CO ₂)	
25	46	0.88	2.53(t,2H,CH ₂ CO ₂); 7.02- 7.38(m,5H,ArH)	
	47	0.87	2.33(t,2H,CH ₂ CO ₂); 7.05- 7.52(m,5H,ArH); 7.21(s,1H,NH)	
	48	0.87	2.33(t,2H,CH ₂ CO ₂); 7.05- 7.52(m,5H,ArH); 7.23(s,1H,NH)	
30	49	0.88	2.33(t,2H,CH ₂ CO ₂); 7.05- 7.54(m,5H,ArH)	
	52	0.90	3.66(s,3H,OMe); 5.83(bs,1H,NH)	
	53		IR. 2140 cm ⁻¹ N ₂	

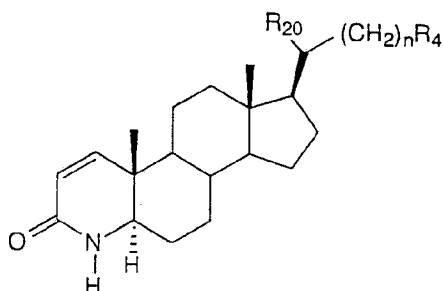
- 39 -

	54	0.90	2.30(t,2H,CH ₂ CO ₂); 3.69(s,3H,OMe)
	55	0.90	7.30(m,1H,ArH); 7.70(m,2H,ArH); 8.62(m,1H,ArH)
5	56	0.92	5.7(bs,1H,NH); 7.31(m,1H,ArH); 7.71(m,2H,ArH); 8.63(m,1H,ArH)
	57	0.90	1.32(d,3H,21-Me); 7.32(m,1H,ArH); 7.72(m,2H,ArH); 8.63(m,1H,ArH)
	59	0.88	0.97(d,3H,21-Me); 3.65(s,3H,OMe)
10	59a	0.88	0.91(d,3H,21-Me); 3.65(s,3H,OMe); 5.77(s,1H,NH)
	62	0.88	2.29(t,2H,CH ₂ CO ₂); 3.66(s,3H,OMe)
	64	0.86	2.33(t,2H,CH ₂ CO ₂)

15

Table 7

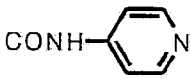
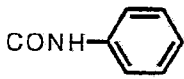
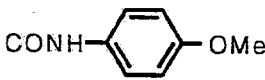
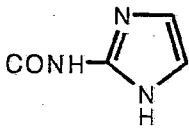
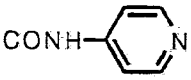
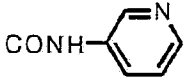
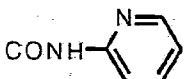
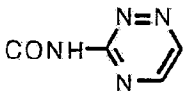
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25

	Compd	R20	n	R4	NMR		Other
					18-Me	19-Me	
	65	Me	0	CO ₂ H			
30	66	Me	0		0.63	0.95	1.33(d,3H,21-Me); 5.81(d,1H,1-H); 6.03(bs,1H,NH); 6.79(d,1H,2-H); 7.34, 7.63, 7.79, 8.64(m,1H ea,ArH)

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5	67	Me	0		0.72	0.94	1.25(d,3H,21-Me); 5.78(d,1H,1-H); 6.79(d,1H,2-H); 7.62(d,2H,ArH); 8.39(d,2H,ArH)
	68	H	1	CO ₂ H	0.57	0.87	5.59(d,1H,1-H); 6.82(d,1H,2-H)
	69	H	1	CO ₂ Me	0.62	0.98	3.66(s,3H,OMe); 5.82(d,1H,1-H); 6.82(d,1H,2-H);
10	70	H	1		0.64	0.98	5.81(d,1H,1-H); 6.82(d,1H,2-H); 7.05- 7.54(m,5H,ArH)
	71	H	1		0.64	0.97	3.79(s,3H,OMe); 5.82(d,1H,1-H); 6.82(d,1H,2-H); 6.86(d,2H,ArH); 7.42(d,2H,ArH)
15	72	H	1		0.62	0.97	5.81(d,1H,1-H); 6.81(d,1H,2-H); 6.78(s,2H,ArH)
	73	H	1		0.64	0.97	5.78(d,1H,1-H); 6.83(d,1H,2-H); 7.58(d,2H,ArH); 8.39(d,2H,ArH)
20	74	H	1		0.63	0.96	5.79(d,1H,1-H); 6.79(d,1H,2-H); 7.28(m,1H,ArH); 8.27(m,2H,ArH); 8.58(s,1H,ArH)
	75	H	1		0.64	0.98	5.81(d,1H,1-H); 6.81(d,1H,2H); 7.04(m,1H,ArH); 7.96(m,1H,ArH); 8.24(m,2H,ArH)
25	76	H	1		0.64	0.97	5.81(d,1H,1-H); 6.80(d,1H,2-H); 8.51(d,1H,ArH); 8.96(d,1H,ArH)
30							

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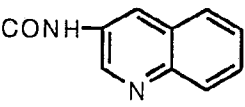
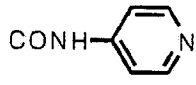
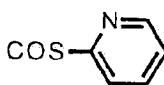
5	77	H	1		0.64	0.95	5.82(d,1H,1-H); 6.86(d,1H,2-H); 7.52- 8.06(m,4H,ArH); 8.71- 8.95(m,2H,ArH)
	78	Me	2				
	79	Me	2				
10	80	Me	2		0.68	0.97	0.95(d,3H,21-Me); 5.46(s,1H,4-NH); 5.81(d,1H,1-H); 6.80(d,1H,2-H); 7.56(d,2H,ArH); 8.13(bs,1H,CONH); 8.46(d,2H,ArH)
	81	Me	2				
	82	Me	2				
20	83	H	1		0.62	0.97	5.80(d,1H,1-H); 6.80(d,1H,2-H); 7.18- 7.78(m,3H,ArH); 8.60(d,1H,ArH)

Table 7, part 2

Compd Mass Spectrum

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	69	m/e359(M+) EI
	70	m/e420(M+) EI
5		
	71	m/e451(M+1) FAB
	72	m/e410(M+) EI
10		
	73	m/e421(M+) EI
	74	m/e422(M+1) FAB
15		
	75	m/e422(M+1) FAB
	76	m/e423(M+) EI
20		
	77	m/e472(M+1) FAB

25

The present invention has the objective of providing suitable topical, oral and parenteral pharmaceutical formulations for use in the novel methods of treatment of the present invention.

30

The compositions containing the compounds of the present invention as the active ingredient for use in the treatment of e.g., benign prostatic hypertrophy, prostatitis, and treatment and prevention of prostatic carcinoma, hyperandrogenic conditions, can be administered in a wide variety of therapeutic dosage forms in conventional vehicles for systemic administration, as, for example,

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by oral administration in the form of tablets, capsules, solutions, or suspensions, or by injection. The daily dosage of the products may be varied over a wide range varying from 0.5 to 1,000 mg per adult human/per day. The compositions are preferably provided in the form of scored tablets containing 0.5, 1.0, 2.5, 5.0, 10.0, 15.0, 25.0, and 50.0 milligrams of the active ingredient for the symptomatic adjustment of the dosage to the patient to be treated. An effective amount of the drug is ordinarily supplied at a dosage level of from about 0.002 mg. to about 50 mg./kg. of body weight per day. Preferably the range is from about 0.01 mg. to 7 mg./kgs. of body weight per day. These dosages are well below the toxic dose of the product. For the treatment of androgenic alopecia, acne vulgaris, seborrhea, female hirsutism, the compounds of the present invention are administered in a pharmaceutical composition comprising the active compound in combination with a pharmacologically acceptable carrier adapted for topical, oral or parenteral administration.

These topical pharmaceutical compositions may be in the form of a cream, ointment, gel or aerosol formulation adapted for application to the skin. These topical pharmaceutical compositions containing the compounds of the present invention ordinarily include about 0.1% to 15%, preferably about 5%, of the active compound, in admixture with about 95% of vehicle.

The compounds of the present invention can be administered in such oral dosage forms as tablets, capsules (each including timed release and sustained release formulations), pills, powders, granules, elixers, tinctures, suspensions, syrups and emulsions. Likewise, they may also be administered in intravenous (both bolus and infusion), intraperitoneal, subcutaneous or intramuscular form, all using forms well known to those of ordinary skill in the pharmaceutical arts. An effective but non-toxic amount of the compound desired can be employed as a 5 α -reductase agent.

The dosage regimen utilizing the compounds of the present invention is selected in accordance with a variety of factors including type, species, age, weight, sex and medical condition of the patient; the

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5 severity of the condition to be treated; the route of administration; the renal and hepatic function of the patient; and the particular compound or salt thereof employed. An ordinarily skilled physician or veterinarian can readily determine and prescribe the effective amount of the drug required to prevent, counter or arrest the progress of the condition. Optimal precision in achieving concentration of drug within the range that yields efficacy without toxicity requires a regimen based on the kinetics of the drug's availability to target sites. This involves a consideration of the distribution, equilibrium, and elimination of a drug.

10 Oral dosages of the present invention, when used for the indicated effects, will range between about Advantageously, compounds of the present invention may be administered in a single daily dose, or the total daily dosage may be administered in divided doses of two, 15 three or four times daily. Furthermore, preferred compounds for the present invention can be administered in intranasal form via topical use of suitable intranasal vehicles, or via transdermal routes, using those forms of transdermal skin patches well known to those of ordinary skill in that art. To be administered in the form of a transdermal delivery 20 system, the dosage administration will, of course, be continuous rather than intermittent throughout the dosage regimen.

In the methods of the present invention, the compounds herein described in detail can form the active ingredient, and are typically administered in admixture with suitable pharmaceutical 25 diluents, excipients or carriers (collectively referred to herein as "carrier" materials) suitably selected with respect to the intended form of administration, that is, oral tablets, capsules, elixirs, syrups and the like, and consistent with conventional pharmaceutical practices.

For instance, for oral administration in the form of a tablet 30 or capsule, the active drug component can be combined with an oral, non-toxic pharmaceutically acceptable inert carrier such as ethanol, glycerol, water and the like. Moreover, when desired or necessary, suitable binders, lubricants, disintegrating agents and coloring agents can also be incorporated into the mixture. Suitable binders include

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starch, gelatin, natural sugars such as glucose or beta-lactose, corn
sweeteners, natural and synthetic gums such as acacia, tragacanth or
sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes
5 and the like. Lubricants used in these dosage forms include sodium
oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium
acetate, sodium chloride and the like. Disintegrators include, without
limitation, starch, methyl cellulose, agar, bentonite, xanthan gum and
the like.

10 The compounds of the present invention can also be
administered in the form of liposome delivery systems, such as small
unilamellar vesicles, large unilamellar vesicles and multilamellar
vesicles. Liposomes can be formed from a variety of phospholipids,
such as cholesterol, stearylamine or phosphatidylcholines.

15 Compounds of the present invention may also be delivered
by the use of monoclonal antibodies as individual carriers to which the
compound molecules are coupled. The compounds of the present
invention may also be coupled with soluble polymers as targetable drug
carriers. Such polymers can include polyvinylpyrrolidone, pyran
20 copolymer, polyhydroxypropyl- methacrylamide-phenol,
polyhydroxyethylaspartamidephenol, or polyethyleneoxidepolylysine
substituted with palmitoyl residues. Furthermore, the compounds of the
present invention may be coupled to a class of biodegradable polymers
useful in achieving controlled release of a drug, for example, polylactic
acid, polyepsilon caprolactone, polyhydroxy butyric acid,
25 polyorthoesters, polyacetals, polydihydropyrans, polycyanoacrylates and
cross-linked or amphipathic block copolymers of hydrogels.

BIOLOGICAL ASSAYS

30 Preparation of Human prostatic and scalp 5 α -reductases.

Samples of human tissue were pulverized using a freezer
mill and homogenized in 40 mM potassium phosphate, pH 6.5, 5 mM
magnesium sulfate, 25 mM potassium chloride, 1 mM phenylmethyl-
sulfonyl fluoride, 1 mM dithiothreitol (DTT) containing 0.25 M sucrose
using a Potter-Elvehjem homogenizer. A crude nuclear pellet was

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prepared by centrifugation of the homogenate at 1,500xg for 15 min. The crude nuclear pellet was washed two times and resuspended in two volumes of buffer. Glycerol was added to the resuspended pellet to a final concentration of 20%. The enzyme suspension was frozen in aliquots at -80°C. The prostatic and scalp reductases were stable for at least 4 months when stored under these conditions.

5 α -reductase assay.

The reaction mixture contained in a final volume of 100 μ l is: 40 mM buffer (human scalp, potassium phosphate, pH 6.5; human prostatic 5 α -reductase, sodium citrate, pH 5.5), 0.3-10 μ M ¹⁴C-T (or ³H-T), 1 mM DTT, and 500 μ M NADPH. Typically, the assay was initiated by the addition of 50-100 μ g prostatic homogenate or 75-200 μ g scalp homogenate and incubated at 37°C. After 10-50 min the reaction was quenched by extraction with 250 μ l of a mixture of 70% cyclohexane: 30% ethyl acetate containing 10 μ g each DHT and T. The aqueous and organic layers were separated by centrifugation at 14,000 rpm in an Eppendorf microfuge. The organic layer was subjected to normal phase HPLC (10 cm Whatman partisil 5 silica column equilibrated in 1 ml/min 70 % cyclohexane: 30 % ethyl acetate; retention times DHT, 6.8-7.2 min; androstanediol, 7.6-8.0; T, 9.1-9.7 min). The HPLC system consisted of a Waters Model 680 Gradient System equipped with a Hitachi Model 655A autosampler, Applied Biosystems Model 757 variable UV detector, and a Radiomatic Model A120 radioactivity analyzer. The conversion of T to DHT was monitored using the radioactivity flow detector by mixing the HPLC effluent with one volume of Flo Scint 1 (Radiomatic). Under the conditions described, the production of DHT was linear for at least 25 min. The only steroids observed with the human prostate and scalp preparations were T, DHT and androstanediol.

Stumptail macaque protocol

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The following protocol is utilized with the stump-tail macaque monkey to demonstrate the effect of compounds of the present invention for promoting hair growth.

5 Twenty-one male stump-tail macaque monkeys of species *Macaca speciosa* are assigned to vehicle control and drug treatment groups on the basis of baseline hair weight data. This assignment procedure is necessary to insure that the average baseline hair growth for each control and experimental group is comparable. The control and
10 drug treatment groups are as follows:

1. Topical 50:30:20 vehicle (N = 6)
2. Oral 5 α -reductase and topical 50:30:20 vehicle (N = 5)
3. Oral placebo (N = 5)
- 15 4. 5 α -reductase in vehicle (N = 5)

The vehicle consists of 50% propylene glycol, 30% ethanol and 20% water. A 100 mM concentration of topical 5 α -reductase is formulated in this vehicle. The same 5 α -reductase is administered as an oral dose of 0.5mg per monkey. Immediately prior to the dosing phase of the study,
20 hair is removed from a 1 inch square area (identified by four tattoos) in the center of the balding scalp. This hair collection is the baseline hair growth determination prior to the beginning of treatment.

Approximately 250 μ L of vehicle and 5 α -reductase in vehicle is prepared and topically administered to the tattooed area of the scalp. The selected
25 5 α -reductase and placebo is ingested by the monkeys at the same time as the topical doses are administered. The monkeys are dosed once per day, seven days per week for twenty weeks.

At four week intervals throughout the dosing phase of the study, each monkey is shaved and the hair is collected and weighed. The
30 body weight data (at baseline and during assay) is analyzed by the nonparametric Wilcoxon rank-sum test. Differences are significant at $p < 0.05$. Hair weight data at each week collection for vehicle, placebo and treatment groups are expressed as the change from baseline.

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Statistical analysis is performed on the rank of the data to show overall differences among groups at each four week collection.

5 While the invention has been described and illustrated with
reference to certain preferred embodiments thereof, those skilled in the
art will appreciate that various changes, modifications and substitutions
can be made therein without departing from the spirit and scope of the
invention. For example, effective dosages other than the preferred
10 dosages as set forth herein above may be applicable as a consequence of
variations in the responsiveness of the mammal being treated for any of
the indications for the compounds of the invention indicated above.
Likewise, the specific pharmacological responses observed may vary
according to and depending upon the particular active compound
15 selected or whether there are present pharmaceutical carriers, as well as
the type of formulation and mode of administration employed, and such
expected variations or differences in the results are contemplated in
accordance with the objects and practices of the present invention. It is
intended, therefore, that the invention be limited only by the scope of
20 the claims which follow and that such claims be interpreted as broadly
as is reasonable.

25

30

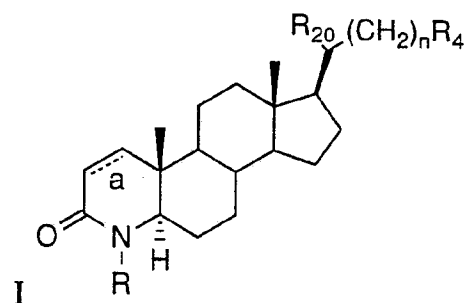
-48A-

WHAT IS CLAIMED IS:

5

1. A compound of the formula:

10



15

wherein:

20

25

30

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dashed line a can represent a double bond when
present;

5 R is selected from hydrogen, methyl or ethyl;

R₂₀ is selected from hydrogen or methyl;

n is an integer from 0 to 10

10

R₄ is selected from:

(a) COR₁, where R₁ is C₆-C₁₀ aryl,
substituted C₆-C₁₀ aryl, and heteroaryl;

15

(b) CONHR₂, where R₂ is substituted phenyl,
heteroaryl, substituted heteroaryl, or C₇ to
C₁₂ cycloalkyl;

20

(c) CO₂R₃, where R₃ is C₆-C₁₀ aryl,
substituted C₆-C₁₀ aryl, or C₇-C₁₂ cycloalkyl;

25

wherein the above aryl or heteroaryl radicals can also be fused with a
benzo or another heteroaryl ring and can further be substituted with
one or more substituents; and the pharmaceutically acceptable salts and
esters thereof.

2. The compound of Claim 1 wherein said aryl
radical is selected from phenyl, benzyl and naphthyl.

30

3. The compound of Claim 1 wherein the heteroaryl
radical is selected from:

pyridyl, pyrrol, thienyl, isothiazolyl, thiazolyl, imidazolyl, tetrazolyl,
pyrazinyl, pyrimidinyl, quinolyl, isoquinolyl, benzothienyl,

- 50 -

isobenzofuryl, pyrazolyl, indolyl, purinyl, carbazolyl, isoxazolyl, triazolyl, furanyl, oxazolyl, or thiadiazolyl.

5 4. The compound of Claim 1 wherein said aryl and heteroaryl substituents are selected from:

hydrogen;

C1-8 straight or branched alkyl;

C2-8 straight or branched alkenyl;

10 C3-8 cycloalkyl;

C2-8 alkynyl;

-CONR⁴R⁵ where R⁴ and R⁵ independently are H, C1-8 alkyl, C3-8 cycloalkyl, C1-4 perhaloalkyl, phenyl, or substituted phenyl, as described below:

15

-COR4;

$-S(O)_nR_4$ where $n=0-2$;

-OCOR4;

-SO₂NR₄R₅;

$$-\text{NR}_4(\text{CO})\text{R}_5;$$

20

-NR₄(CO)NHR₅;

-NHSO₂R₄;

-OR4;

-NR4R5;

CN;

25

NO2;

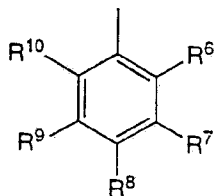
halo;

perhalo C1-C4alkyl;

$$-\text{CO}_2\text{R}_4;$$

phenyl or substituted phenyl of the formula:

30



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where R6-R10 independently represent one or more of the substituents as defined above.

5 5. The compound of claim 1 being the following:

21-benzoyl-4-aza-5a-pregnan-3-one,

21-benzoyl-4-methyl-4-aza-5a-pregnan-3-one,

10 21-(2-methoxybenzoyl)-4-aza-5a-pregn-1-ene-3- one,

4-methyl-21-(2-trifluoromethylbenzoyl)-4-aza- 5a-
pregn-1-en-3-one,

15 20-benzoyl-4-methyl-4-aza-5a-pregnan-3-one,

23-(2-fluorobenzoyl)-4-methyl-24-nor-4-aza-5a- cholane-
3,23-dione,

20 23-(3-pyridyl)-24-nor-4-aza-5a-chol-1-ene- 3,23-
dione,

4-methyl-24-(2-pyridyl)-21-nor-4-aza-5a- cholane-3,24-
dione,

25 17b-[5-(3-chlorobenzoyl)pentyl]-4-aza-5a- androstan-3-
one,

30 17b-(6-benzoylhexyl)-4-methyl-4-aza-5a- androstan-3-
one,

17b-(10-benzoyldecyl)-4-aza-5a-androst-1-en-3- one.

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- dione, 4-methyl-21-(2-thienyl)-4-aza-5a-pregnane- 3,21-
- 5 24-(2-pyrazinyl)-4-aza-5a-chol-1-ene-3,24- dione,
- 4-ethyl-(2,6-dimethoxybenzoyl)-4-aza-5a- pregnan-3-one,
- 10 N-(4-acetylphenyl)-4-methyl-3-oxo-4-aza-5a-
pregnane-21-carboxamide,
- N-(4-acetylphenyl)-3-oxo-4-aza-5a-pregnane-21-
carboxamide,
- 15 4-methyl-3-oxo-N-(4-pyridyl)-4-aza-5a-preg-
nane-21-carboxamide,
- 3-oxo-N-(4-pyridyl)-4-aza-5a-pregnane-21-car-
boxamide,
- 20 N-(2-adamantyl)-3-oxo-4-aza-5a-pregnane-21-
carboxamide,
- N-(2-adamantyl)-4-methyl-3-oxo-4-aza-5a- pregnane-21-
carboxamide,
- 25 3-oxo-N-(4-pyridyl)-4-aza-5a-pregnan-21-amide,
- 4-methyl-3-oxo-4-(4-pyridyl)-4-aza-21-nor-5a- cholan-
24-amide,
- 30 4-methyl-3-oxo-N-(3-pyridyl)-4-aza-5a- pregnane-21-
carboxamide,
- 4-methyl-3-oxo-N-(2-pyridyl)-4-aza-5a-

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pregnane-21-carboxamide,

5 N-(1-adamantyl)-4-methyl-3-oxo-4-aza-5a- pregnane-
20(S)-carboxamide,

N-(4-acetylphenyl)-4-methyl-3-oxo-4-aza-5a-
pregnane-20(S)-carboxamide,

10 N-(4-chlorophenyl)-4-methyl-3-oxo-4-aza-5a- cholan-
24-amide,

N-(4-acetylphenyl)-4-methyl-3-oxo-4-aza-5a-
cholan-24-amide,

15 3-oxo-N-(4-trifluoromethylphenyl)-4-aza-5a-
cholan-24-amide,

20 4-methyl-3-oxo-N-(4-pyridyl)-24-nor-4-aza-5a-
cholan-24-amide,

N-(1-adamantyl)-11-(4-methyl-3-oxo-4-aza-5a-
androstan-17 β -yl)undecanamide,

25 N-(2-pyridyl)-6-(4-methyl-3-oxo-4-aza-5a-
androstan-17 β -yl)hexanamide,

N-(3-pyridyl)-5-(3-oxo-4-aza-5a-androst-1-en- 17 β -
yl)pentanamide,

30 N-(2-thienyl)-7-(4-methyl-3-oxo-4-aza-5a- androstan-17 β -
yl)heptanamide,

3-oxo-N-(2-pyrazinyl)-4-aza-5a-pregnan-21- amide.

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- 4-methyl-3-oxo-N-(2-t-butylphenyl)-4-aza-5a- cholane-
24-carboxamide,
- 5 4-methyl-3-oxo-N-(2-cyanophenyl)-4-aza-chol-1-
ene-24-carboxamide,
- N-(2-bicyclo[2.2.2]octyl)-9-(3-oxo-4- aza-5a-
androstan-17 β -yl)nonanamide,
- 10 1-adamantyl 4-methyl-3-oxo-4-aza-5a-pregnane- 20(S)-
carboxylate,
- phenyl 4-methyl-3-oxo-4-oxo-4-aza-5a-pregnane-
20(S)-carboxylate,
- 15 2-(t-butyl)phenyl 4-methyl-3-oxo-4-aza-5a-
pregnane-21-carboxylate,
- 2-methoxyphenyl 4-methyl-3-oxo-4-aza-5a-preg
nane-21-carboxylate,
- 20 phenyl 3-oxo-4-aza-5a-pregnane-21-carboxylate,
- phenyl 4-methyl-3-oxo-4-aza-5a-pregnane-21-
carboxylate,
- 25 phenyl 5-(4-methyl-3-oxo-4-aza-5a-androstan- 17 β -
yl)pentanoate,
- 30 2-(t-butyl)phenyl 3-oxo-4-aza-5a-pregnan-21- oate,
- 2,6-dimethoxyphenyl 3-oxo-4-aza-5a-pregn-1-en-
21-oate.

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2-adamantyl 8-(4-methyl-3-oxo-4-aza-5a-androstan-17 β -yl)octanoate,

5 2,6-dimethylphenyl 3-oxo-4-aza-5a-pregn-1-en-21-oate,

2,6-dichlorophenyl 4-methyl-3-oxo-4-aza-5a-pregn-1-en-21-ate,

10 phenyl 10-(4-methyl-3-oxo-4-aza-5a-androstan-17 β -yl)decanoate,

6. A method for treating the hyperandrogenic conditions of acne, androgenic alopecia, male pattern baldness, female hirsutism, benign prostatic hyperplasia, prostatitis, treatment and
15 prevention of prostatic cancer, comprising administering to a patient in need of such treatment a therapeutically effective amount of the compound of Claim 1.

20 7. The method of Claim 6 wherein said compound is a 5 α -reductase 1 inhibitor.

8. The method of Claim 6 wherein said compound is a 5 α -reductase 2 inhibitor.

25 9. The method of Claim 6 wherein said compound is a dual inhibitor for both 5 α -reductase 1 and 2.

30 10. A pharmaceutical composition comprising a therapeutically effective amount of the compound of Claim 1 in a pharmaceutically acceptable vehicle therefor.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US93/04631

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) :A61K 31/495; C07D 241/12

US CL :544/406; 514/253

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : A61K 470, 435; 546/77, 78; 514/284

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS Online Structure Search

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	Jour. Clinical Endoc. and Metabl., Vol. 74 1992. Diani et al. "HAIR GROWTH EFFECTS OF ORAL ADMINISTRATION OF FINA STERIDE, A STEROID 5 α Reductase Inhibitor, Alone and in combination with Topical Minoxidil in the Balding stump-tail macaque, pages 345-350. See page 3456, para. bridging cols. 1-2, last 3 lines.	3-10

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

<p>* Special categories of cited documents:</p>	
A document defining the general state of the art which is not considered to be part of particular relevance	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
E earlier document published on or after the international filing date	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
O document referring to an oral disclosure, use, exhibition or other means	*&* document member of the same patent family
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

28 JULY 1993

Date of mailing of the international search report

21 SEP 1993

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/US93/04631

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	J. ORG. CHEM., Vol. 46, 1981, Back "OXIDATION OF AZASTEROID LACTAMS AND ALCOHOLS WITH BENZENE SELENIC ANHYDRIDE" pages 1442-6	3-10
A	Jour. Med. Chem., Vol. 27, 1984, Rasmusson et al. "AZASTEROIDS AS INHIBITORS OF RAT PROSTATIC 5 α REDUCTASE" pages 1690-1701. See entire document.	1 and 3-10
X	Jour. med. Chem., Vol. 29, 1986 Rasmusson et al.	<u>1, 3-6</u>
Y	"AZASTEROIDS, STRUCTURE-ACTIVITY RELATIONSHIPS FOR INHIBITION OF 5 α -REDUCTASE and of ANDROGEN RECEPTOR BINDING" pages 2298-2315. See page 2300 compounds 10ai, 10ac, 9av, 10av, p. 239. Scheme 3	1, 3-10
X	EP, B, 0,200,859 (Cainelli et al.) 12 November 1986, See page 2, line 15 - page 3, line 4, page 5, lines 7 to 11.	<u>1, 4, 6-10</u> 1, 3-10
A	US, A, 4,377,584 (RASMUSSON et al.) 22 March 1983. See entire document.	1 and 3-10
A	us, a, 4,760,071 (RASMUSSON et al.) 26 July 1988. See entire document.	1 and 3-10
A	US, A, 4,882,319 (HOLT et al.) 21 November 1989. See entire document.	1 and 3-10
A	US, A, 5,049,562 (RASMUSSON et al.) 17 September 1991. See entire document.	1 and 3-10
A	US, A, 5,116,983 (BHAHACHARYA et al.) 26 May 1992. See entire document.	1 and 3-10
A	US, A, 5,110,939 (HOLT et al.) 05 May 1992. See entire document.	1 and 3-10
A	Back et al. "N-CHLOROAZASTEROIDS: A NOVEL CLASS OF REACTIVE STEROID ANALOGUES, PREPARATION, REACTION WITH THIOLS AND PHOTOCHEMICAL CONVERSION TO TO ELECTROPHILIC N-ACYL IMINES" pages 1904-10, See entire document.	1 and 3-10
Y	US, A, 4,859,681 (RASMUSSON et al.) 22 August 1989. See abstract.	1-6

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US93/04631

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING

This ISA found multiple inventions as follows:

- I. Claims 1 and 3-10 (part of each), drawn to pyrazines, classified in class 544, subclass 336 plus and 514/253.
- II. Claims 1 and 3-10 (part of each) and 2, drawn to compounds with no heteroaryls, classified in class 546, subclass 77 and 514/284, and drawn to heteroaryls (5-member or pyridyl) not in I or III-VI, classified in class 546, subclass 77 and 514/284.
- III. Claims 1, 3-4 and 6-10 (part of each), drawn to pyrimidines, classified in class 544, subclasses 242 plus and 514/269 plus.
- IV. Claims 1, 3-4 and 6-10 (part of each), drawn to purines, classified in Class 544, subclasses 264 plus and 514/261 plus.
- V. Claims 1, 4 and 6-10 (part of each) drawn to triazines, classified in Class 544, subclasses 180 plus and 514/241 and 43.
- VI. Claims 1, 4 and 6-10 (part of each), drawn to 6 or 7 membered multi N heterocyclics or N plus another hetero 6 or 7 membered heterocyclic not provided above classified in Class 544, subclasses various and 514/various.

Claims, groups lacks a common "a special technical feature, PCT Rule 13.2, not also shared with Carnelli EP and Rasmusson 1986, the prior art. Lack of Unity of invention is not precluded by including in a single claim PCT Rule 13.3.